

Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choepel, Y., Collangelo, M., Collins, S., Collimore, A., Cook, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S., Faro, S., Ferreira, P., Fitzgerald, M., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRoque, K., Lamazares, R., Landers, T., Lehoczyk, J., Levine, R., Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupback, R., Seaman, S., Severy, P., Spencer, B., Strange-Thomann, N., Stojanovic, N., Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
Submitted (09-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
3 (bases 1 to 194420)

Bitren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N., Anderson, M., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Choepel, Y., Collimore, A., Cook, A., Cooke, P., Corum, B., DeArellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, X., Lui, A., Mabbitt, R., MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schupback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Strange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
Submitted (18-DEC-2003) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA  
On Dec 18, 2003 this sequence version replaced gi:38259237.  
All repeats were identified using RepeatMasker:  
Smit, A.F.A. & Green, P. (1996-1997)  
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center  
Center: Whitehead Institute/ MIT Center for Genome Research  
Center code: WIBR  
Web site: <http://www-seq.wi.mit.edu>  
Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)  
----- Project Information  
Center project name: L23552  
Center clone name: 218\_L\_23

\* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. Gaps between the contigs are represented as runs of N. The order of the pieces is believed to be correct as given, however the sizes of the gaps between them are based on estimates that have been provided by the submittor.  
\* This sequence will be replaced  
\* by the finished sequence as soon as it is available and  
\* the accession number will be preserved.  
\* 1 115229: contig of 115229 bp in length  
\* 115230 115329: gap of 100 bp  
\* 115330 194420: contig of 79091 bp in length.

FEATURES  
source

Location/Qualifiers  
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/chromosome="15"  
/map="15"  
/clone="RP23-218L23"  
/clone\_lib="RPCI-23 Female Mouse BAC"

## ORIGIN

Alignment Scores:  
Pred. No.: 374 Length: 194420  
Score: 58.00 Matches: 11  
Percent Similarity: 70.59% Conservative: 1  
Best Local Similarity: 64.71% Mismatches: 5  
Query Match: 58.59% Indels: 0  
DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC124133 (1-194420)

Oy 1 MetaAlaHisSerValLeuSerPheIleuThrProTyAlaLeu 17

Db 29103 ATGAACCTTCACAAAGTCCTTCTATCTCTCTGGACACCATAGTCTC 29053

## RESULT 8

AC096099/c

## LOCUS

AC096099 244669 bp DNA linear HTG 13-NOV-2002

Rattus norvegicus clone CH230-27M23, WORKING DRAFT SEQUENCE.

## DEFINITION

AC096099

AC096099.6 GI:24941648

HTG; HTGS PHASE2; HTGS DRAFT; HTGS\_FULLTOP.

## KEYWORDS

Rattus norvegicus (Norway rat)

## SOURCE

Rattus norvegicus

## ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

## REFERENCE

1 (bases 1 to 244669)

## AUTHORS

Muzny, D., Marie, Metzker, M., Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Aryalabechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Bissio, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flag, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hamilton, K., Harveys, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S., Hodgson, A., Hogues, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpach, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensu, L., Louised, H., Lozano, R., Lu, X., Ma, J., Maheshwari, M., Mahindaratne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwakaeme, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Pu, L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R.,



\* be preserved.

1 1085: contig of 1085 bp in length  
 \* 1086 1185: gap of unknown length  
 \* 1186 11944: contig of 10759 bp in length  
 \* 11945 12044: gap of unknown length  
 \* 12045 30225: contig of 18181 bp in length  
 \* 30226 30325: gap of unknown length  
 \* 30326 63384: contig of 33059 bp in length  
 \* 63385 117854: contig of 54370 bp in length  
 \* 117855 166434: gap of unknown length  
 \* 117955 166435: contig of 48480 bp in length  
 \* 166435 245830: gap of unknown length  
 \* 166535 245830: contig of 79296 bp in length.

## FEATURES

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## ORIGIN

Alignment Scores:  
 Pred. No.: 488 Length: 245830  
 Score: 58.00 Matches: 11  
 Percent Similarity: 70.59% Conservative: 1  
 Best Local Similarity: 64.71% Mismatches: 5  
 Query Match: 58.59% Indels: 0  
 DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC140313 (1-245830)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeu 17  
 Db 14529 ATGAAGACTTCACAAAGTCCTTCTATCCTCTGTGGACCCCATAGTC 14479

RESULT 10  
 AC013568/c 156471 bp DNA linear HTG 26-MAY-2000  
 LOCUS  
 DEFINITION Homo sapiens clone RP11-1B9, WORKING DRAFT SEQUENCE, 10 unordered  
 pieces.

ACCESSION AC013568  
 VERSION AC013568.3 GI:7107832  
 KEYWORDS HTG; HTGS PHASE1; HTGS\_DRAFT.  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 156471)  
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,  
 Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,  
 Brown,A., Castle,A., Colangelo,M., Collins,S., Collymore,A.,  
 Cooke,P., DeArallano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,  
 Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,

Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,  
 Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,  
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 McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,  
 Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,  
 Peterson,K., Follara,V., Riley,R., Roy,A., Santos,R., Severy,P.,  
 Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,  
 Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,  
 Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

## TITLE

## JOURNAL

Submitted (13-NOV-1999) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 On Feb 28, 2000 this sequence version replaced gi:6514007.

## COMMENT

All repeats were identified using RepeatMasker:  
 Smit, A.F.A. & Green, P. (1996-1997)  
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research  
 Center code: WIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence\_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L2468

Center clone name: 1\_B9

----- Summary Statistics

Sequencing vector: M13; M7815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 136271 bases at least Q40

Consensus quality: 149734 bases at least Q30

Consensus quality: 153894 bases at least Q20

Insert size: 170000; agarose-fp

Insert size: 155571; sum-of-contigs

Quality coverage: 5.4 in Q20 bases; agarose-fp

Quality coverage: 5.9 in Q20 bases; sum-of-contigs

\* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 10 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

1 1916: contig of 1916 bp in length  
 \* 1917 2016: gap of 100 bp  
 \* 2017 4993: contig of 2977 bp in length  
 \* 4994 5094: gap of 100 bp  
 \* 5094 10286: contig of 5193 bp in length  
 \* 10287 10386: gap of 100 bp  
 \* 10387 15563: contig of 5177 bp in length  
 \* 15564 15663: gap of 100 bp  
 \* 15664 27659: contig of 11996 bp in length  
 \* 27660 27759: gap of 100 bp  
 \* 27760 41302: contig of 13543 bp in length  
 \* 41303 41402: gap of 100 bp  
 \* 41403 54013: contig of 12611 bp in length  
 \* 54014 54113: gap of 100 bp  
 \* 54114 75521: contig of 21408 bp in length  
 \* 75522 75621: gap of 100 bp  
 \* 75622 109229: contig of 33508 bp in length  
 \* 109130 109229: gap of 100 bp  
 \* 109230 156471: contig of 47242 bp in length.

## FEATURES

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 /note="assembly\_fragment"  
 misc\_feature 2017. 4993





JOURNAL	Submitted (01-MAR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	source	1. .81715
REFERENCE	3 (bases 1 to 81715)		/organism="Homo sapiens"
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzgerald,M., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A., Karatas,A., Kellis,C., LaRocque,K., Lamazares,R., Landers,T., Lenoczky,J., Levine,R., Lindblad-Toh,K., Liu,G., MacLean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	repeat_region	/db_type="genomic DNA"
	Direct Submission	repeat_region	/db_xref="taxon:9606"
TITLE	Submitted (21-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	repeat_region	/chromosome="8"
JOURNAL	4 (bases 1 to 81715)	repeat_region	/clone="RP11-347D13"
REFERENCE	Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzgerald,M., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A., Karatas,A., Kellis,C., LaRocque,K., Lamazares,R., Landers,T., Lenoczky,J., Levine,R., Lindblad-Toh,K., Liu,G., MacLean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.	repeat_region	/complement(1..37)
AUTHORS	Direct Submission	repeat_region	/rpt_family="Alu"
	Submitted (25-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA	repeat_region	/complement(40..342)
JOURNAL	Research, 320 Charles Street, Cambridge, MA 02141, USA	repeat_region	/rpt_family="AluSg"
REFERENCE	On Jun 25, 2002 this sequence version replaced gi:21536046.	repeat_region	/complement(810..867)
COMMENT	All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) <a href="http://ftp.genome.washington.edu/RM/RepeatMasker.html">http://ftp.genome.washington.edu/RM/RepeatMasker.html</a>	repeat_region	/rpt_family="MER5B"
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	Center: Whitehead Institute/ MIT Center for Genome Research	repeat_region	/rpt_family="AluY"
	Center code: WIBR	repeat_region	1753..2182
	Web site: <a href="http://www-seq.wi.mit.edu">http://www-seq.wi.mit.edu</a>	repeat_region	/rpt_family="MLT1D"
	Contact: <a href="mailto:sequence_submissions@genome.wi.mit.edu">sequence_submissions@genome.wi.mit.edu</a>	repeat_region	3424..3730
	----- Project Information	repeat_region	/rpt_family="AluSx"
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	Center Clone name: 347_D_13	repeat_region	/rpt_family="MIR"
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		repeat_region	/rpt_family="L1MB5"
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Only the last 81.7 kb of this clone are being submitted.  
The remainder overlaps accession number AC079089 [WICR project L10641].

Location/Qualifiers

```

REFERENCE
AUTHORS
3 (bases 1 to 165127)
Barren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Miengwa,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (20-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 165127)
Barren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., FitzGerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Miengwa,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (20-SEP-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Sep 20, 2002 this sequence version replaced gi:22325339.
All repeats were identified using RepeatMasker:
Smit,A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
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Center: Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
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Center project name: L10484
Center clone name: 102_F_4
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/db_xref="taxon:9606"
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/map="8"
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/clone_lib="RP11-11 Human Male BAC"
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rpt_family="L2"
1797..1803
/note="<30 qual SNGL region"
1821..1825

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repeat_region /rpt_family="CAAA)n"
repeat_region 2805..3112
repeat_region /rpt_family="AluSg"
repeat_region complement(3673..3776)
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repeat_region /rpt_family="MER82"
repeat_region complement(5020..5063)
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repeat_region complement(5070..5378)
repeat_region /rpt_family="L1MC4a"
repeat_region 5392..5444
repeat_region /rpt_family="(TCTA)n"
repeat_region complement(5446..5567)
repeat_region /rpt_family="L1MC4"
repeat_region 5935..5955
repeat_region /rpt_family="AT_rich"
repeat_region 6175..6258
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repeat_region /rpt_family="L1ME"
repeat_region 9147..9309
repeat_region /rpt_family="MER103"
repeat_region complement(10056..10952)
repeat_region /rpt_family="L1PAL0"
repeat_region complement(12015..12381)
repeat_region /rpt_family="THEIC"
repeat_region 12402..12570
repeat_region /rpt_family="MIR"
repeat_region complement(12765..12978)
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repeat_region complement(13005..13247)
repeat_region /rpt_family="AluSc"
repeat_region complement(13285..13325)
repeat_region /rpt_family="AluSc"
repeat_region complement(14540..14604)
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repeat_region 14605..14902
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repeat_region 21832..22130
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repeat_region 22358..22661
repeat_region /rpt_family="AluJo"
repeat_region 23062..23127
repeat_region /rpt_family="(CA)n"
repeat_region 23297..23425
repeat_region /rpt_family="MIR"
repeat_region 23516..23604
repeat_region /rpt_family="MIR"
repeat_region complement(23704..24014)
repeat_region /rpt_family="AluSx"
repeat_region 24368..24523
repeat_region /rpt_family="MIR3"
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repeat_region complement(24787..24892)
repeat_region /rpt_family="MLT1J"

Alignment Scores:
Pred. No.: 717 Length: 165127
Score: 56.00 Matches: 9
Percent Similarity: 78.57% Conservative: 2
Best Local Similarity: 64.29% Mismatches: 3
Query Match: 56.57% Indels: 0
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AC091047 (1-165127)
QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTriThrPro 14
Db 33611 ATGCCACAACTCGATTGACGTTTATTCTGACCCCA 33570

RESULT 14
AC140381 233717 bp DNA linear HTG 23-FEB-2003
LOCUS Mus musculus chromosome UNK clone RP23-176M4, WORKING DRAFT
DEFINITION SEQUENCE, 18 unordered pieces.
ACCESSION AC140381
VERSION AC140381.1 GI:28475640
KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS_FULLTOP.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 233717)
AUTHORS McPherson,J.D. and Waterston,R.H.
TITLE The sequence of Mus musculus clone
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 233717)
AUTHORS McPherson,J.D. and Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (23-FEB-2003) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA

COMMENT
----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc/index.shtml
Contact: submissions@wustl.wustl.edu
----- Project Information -----
Center project name: M_BA0176M04
----- Summary Statistics -----
Sequencing vector: M13, 0%
Sequencing vector: plasmid, 100%
Chemistry: Dye-primer ET; 0% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 226925 bases at least Q40
Consensus quality: 228328 bases at least Q30
Consensus quality: 229211 bases at least Q20
* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
```





RP11-117113 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.  
The true left end of clone RP11-117113 is at 1 in this sequence.  
The true left end of clone RP11-366017 is at 98459 in this sequence.

## FEATURES

Source Location/Qualifiers

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4303..4506
/note="L1MA9 repeat: matches 6070..6270 of consensus"
repeat_region
4536..4844
/note="AluSg repeat: matches 1..305 of consensus"
repeat_region
5066..5362
/note="AluSx repeat: matches 1..298 of consensus"
repeat_region
5544..5674
/note="L2 repeat: matches 2619..2748 of consensus"
repeat_region
7241..8826
/note="L1PB1 repeat: matches 4551..6148 of consensus"
repeat_region
8828..9507
/note="L1M1 repeat: matches 2438..2734 of consensus"
repeat_region
9596..10327
/note="L1PBa repeat: matches -237..1442 of consensus"
repeat_region
10350..11388
/note="L1PBa repeat: matches -1537..-412 of consensus"
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12251..12425
/note="match: STS: Em:G15057"
repeat_region
12414..12652
/note="MER57-internal repeat: matches 7155..7387 of consensus"
repeat_region
12876..13029
/note="L1M1H repeat: matches 178..340 of consensus"
repeat_region
13220..13577
/note="L1M1H repeat: matches 43..410 of consensus"
repeat_region
13578..13658
/note="L2 repeat: matches 2420..2496 of consensus"
repeat_region
13659..14025
/note="THE1C repeat: matches 1..369 of consensus"
repeat_region
14026..14254
/note="L2 repeat: matches 2201..2420 of consensus"
repeat_region
14255..14532
/note="AluJ0 repeat: matches 1..280 of consensus"
repeat_region
14533..14844
/note="L2 repeat: matches 1862..2201 of consensus"
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15852..16106
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16209..16274
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16270..16309
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16351..16819
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17215..17470
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17722..17851
/note="MIR repeat: matches 61..191 of consensus"
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18490..18521
/note="16 copies 2 mer tg 93% conserved"
repeat_region
19327..19402
/note="L1M1F repeat: matches 89..161 of consensus"
repeat_region
19430..19465
/note="18 copies 2 mer tc 86% conserved"
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19536..19782
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20422..20618
/note="L2 repeat: matches 2554..2747 of consensus"
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/note="AluSp/q repeat: matches 187..303 of consensus"
21647..21868
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22326..22377
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24000..24027
/note="14 copies 2 mer tg 89% conserved"
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25652..25812
/note="MER44A repeat: matches 5..170 of consensus"
25813..26530
/note="trigger3(Golem) repeat: matches 2277..3028 of consensus"
26524..26594
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26655..26806
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26817..27149
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29151..29504
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32372..32588
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33053..33329
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33807..33871
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34562..34746
/note="MERSA repeat: matches 1..188 of consensus"
35475..35635
/note="MIR repeat: matches 78..230 of consensus"
35840..36257
/note="L2 repeat: matches 1450..1892 of consensus"
38733..39038
/note="L1M1 repeat: matches 142..468 of consensus"
39053..39147
/note="MER86 repeat: matches 88..182 of consensus"
39567..39723
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40423..40607
/note="L1M4 repeat: matches 2680..2886 of consensus"
40608..41117
/note="LTR40a repeat: matches 1..519 of consensus"
41118..41187
/note="L1M4 repeat: matches 2886..2950 of consensus"
41302..41405
/note="52 copies 2 mer aa 60% conserved"
42443..42709
/note="L2 repeat: matches 2457..2710 of consensus"
43036..43484
/note="L1M1C repeat: matches 7..466 of consensus"
44299..44582
/note="AluSx repeat: matches 1..296 of consensus"
46846..46936
/note="MER91A repeat: matches 17..113 of consensus"

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repeat_region 48664..49299
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repeat_region 49298..49537
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repeat_region 49608..49961
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repeat_region 51260..51623
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## Alignment Scores:

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Pred. No.:      607      Length:      98558
Score:          55.00      Matches:      9
Percent Similarity: 100.00%      Conservative: 5
Best Local Similarity: 64.29%      Mismatches: 0
Query Match:      55.56%      Indels: 0
DB:              9      Gaps: 0

```

US-10-799-747-116 (1-20) x AL137138 (1-98558)

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QY      2 AlalahisservalleuSerPheLeuLeuTrpThrProTyr I5
Db      25528 GCAAGCCACACAGTCATCTCTTCTGCTATATACCTTC 25487

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Search completed: July 21, 2004, 02:13:58  
Job time : 1837 secs

**This Page Blank (uspto)**

GenCore version 5.1.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 00:02:30 ; Search time 348 Seconds  
(without alignments)  
244.149 Million cell updates/sec

Title: US-10-799-747-116  
Perfect score: 99  
Sequence: 1 MAHSVLSFLWTPYALKSX 20

Scoring table:  
BLOSUM62  
Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 3373863 seqs, 2124099041 residues

Total number of hits satisfying chosen parameters: 6747726

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000  
Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Command line parameters:

-MODEL=frame+p2n.model -DEV=xlp  
-O=/cpn2.1/USPTO.spool.p/US10799747/runat.19072004.161415.21434/app.query.fasta\_1.199  
-DB=N\_Geneseq\_29Jan04 -QFMT=fastap -SUFFIX=ring -MINMATCH=0.1 -LOOPEL=0  
-LOOPEXT=0 -UNITIS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi  
-LIST=45 -DOCALLIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15  
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10799747 @CGN 1 1 708 @runat.19072004.161415.21434 -NCPU=6 -ICPU=3  
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPELOCK=100 -LONGLOG  
-DEV\_TIMEOUT=120 -WARN\_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6  
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : N\_Geneseq\_29Jan04.\*

- 1: Geneseqn1980s.\*
- 2: Geneseqn1990s.\*
- 3: Geneseqn2000s.\*
- 4: Geneseqn2001as.\*
- 5: Geneseqn2001bs.\*
- 6: Geneseqn2002s.\*
- 7: Geneseqn2003as.\*
- 8: Geneseqn2003bs.\*
- 9: Geneseqn2003cs.\*
- 10: Geneseqn2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	98	99.0	1434	2 AAX37452	Human sec
2	98	99.0	1434	7 ADA39771	Human sec
3	98	99.0	1434	7 ACC50424	Human sec
c	56	56.6	8417	4 AAK89495	Human dig
5	56	56.6	8417	4 AAS32681	Human gen
6	53	53.5	5760	4 AAS32472	Human gen
7	53	53.5	5763	4 AAS32471	Human gen
8	52	52.5	2877	7 ABZ36220	Human sec

c	9	52	52.5	36785	4	AAX82208	Human imm
c	10	51	51.5	400	5	AAF64369	Novel hum
c	11	51	51.5	505	4	AAH03292	Human cDN
c	12	51	51.5	708	5	AAF64473	Novel hum
c	13	51	51.5	1544	2	AAT86839	cDNA enco
c	14	51	51.5	1568	4	AAH15239	Human cDN
c	15	51	51.5	2500	4	AAH77480	Mandrill
c	16	51	51.5	9641	7	AAH77486	Mandrill
c	17	50	50.5	620	7	ABT40513	Toxicity
c	18	50	50.5	620	9	ADB50026	Primary r
c	19	49	49.5	6708	9	ADC56730	Thale cre
c	20	49	49.5	6708	9	ADC56727	Open read
c	21	49	49.5	7149	9	ADC56728	Thale cre
c	22	49	49.5	11000	9	ADC56726	Thale cre
c	23	49	49.5	32768	2	AAX13065	Enterococ
c	24	49	49.5	32768	6	ABS98860	Enterococ
c	25	48	48.5	207	7	ABX88609	Corn ear-
c	26	48	48.5	290	2	AAV88159	EST clone
c	27	48	48.5	375	4	AAS39268	Novel hum
c	28	48	48.5	404	3	AAF11614	Aspergill
c	29	48	48.5	481	6	ABN64079	Human can
c	30	48	48.5	485	5	ABV45709	Human pro
c	31	48	48.5	495	6	ABO57387	Human col
c	32	48	48.5	612	6	ABY07938	Human lun
c	33	48	48.5	648	5	ABV15807	Human pro
c	34	48	48.5	675	5	ABV45609	Human pro
c	35	48	48.5	804	4	AAH32323	Human olf
c	36	48	48.5	939	7	ABZ77940	Human G p
c	37	48	48.5	940	9	ADC86574	Human GPC
c	38	48	48.5	960	4	AAH12959	Human G-p
c	39	48	48.5	960	5	AAS42282	Human cDN
c	40	48	48.5	960	6	ABZ43079	Human GPC
c	41	48	48.5	960	6	ABK68434	Human DNA
c	42	48	48.5	961	6	ABK37568	DNA encod
c	43	48	48.5	961	6	ABD28710	Human G-p
c	44	48	48.5	970	6	ABS58783	Human G-p
c	45	48	48.5	970	6	AAD28709	Human G-p

#### ALIGNMENTS

RESULT 1  
AAX37452  
ID AAX37452 standard; cDNA; 1434 BP.  
XX  
AC AAX37452;  
XX  
DT 06-JUL-1999 (first entry)  
XX  
DE Human secreted protein cDNA fragment containing gene 2.  
XX

Human; secreted protein; treatment; prevention; protein therapy; AIDS;  
gene therapy; diagnosis; cancer; tumour; neurodegenerative disorder;  
developmental abnormality; fetal deficiency; blood disorder; leukemia;  
immune system disease; autoimmune disease; hepatic disease; lymphoma;  
renal disease; inflammation; allergy; Alzheimer's disease; schizophrenia;  
cognitive disorder; prostate disease; skeletal; cardiac; muscle disorder;  
pulmonary disorder; transplant rejection; osteoclast; osteoporosis;  
arthritis; malignancy; digestive; infection; ss.

OS Homo sapiens.

XX  
XX WO9918208-A1.  
PN  
XX  
PD 15-APR-1999.  
XX  
XX 01-OCT-1998; 98WO-US020775.  
PF  
XX  
XX 02-OCT-1997; 97US-0060833P.  
PR  
XX 02-OCT-1997; 97US-0060833P.  
PR  
XX 02-OCT-1997; 97US-0060837P.  
PR  
XX 02-OCT-1997; 97US-0060838P.  
PR  
XX 02-OCT-1997; 97US-0060838P.  
PR

PR 02-OCT-1997; 97US-0060843P.  
 PR 02-OCT-1997; 97US-0060862P.  
 PR 02-OCT-1997; 97US-0060866P.  
 PR 02-OCT-1997; 97US-0060874P.  
 PR 02-OCT-1997; 97US-0060880P.  
 PR 02-OCT-1997; 97US-0060884P.  
 XX  
 PA (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 XX Duan DR, Florence KA, Rosen CA, Ruben SM, Greene JM, Young P;  
 PI Ferrie AM, Yu G, Janat F, Ni J, Carter KC, Endress GA, Feng P;  
 PI Lafleur DW, Shi Y;  
 XX  
 DR WPI; 1999-264022/22.  
 DR P-PSDB; AAY07853.  
 XX  
 XX New isolated human genes and the secreted polypeptides they encode.  
 PT  
 XX  
 XX Claim 1a; Page 228; 368pp; English.  
 XX  
 CC This invention describes novel isolated human genes and the secreted  
 CC proteins they encode. The products of the invention are useful for  
 CC preventing, treating or ameliorating medical conditions, e.g. by protein  
 CC or gene therapy. Also pathological conditions can be diagnosed by  
 CC determining the amount of the new polypeptides in a sample or by  
 CC determining the presence of mutations in the new polynucleotides.  
 CC Specific uses are described for each of the 101 polynucleotides, based on  
 CC which tissues they are most highly expressed in, and include developing  
 CC products for the diagnosis or treatment of cancer, tumours,  
 CC neurodegenerative disorders, developmental abnormalities and fetal  
 CC deficiencies, blood disorders, leukemias, diseases of the immune system,  
 CC autoimmune diseases, hepatic and renal disease, lymphomas, inflammation,  
 CC allergies, Alzheimer's and cognitive disorders, schizophrenia, prostate  
 CC disease, skeletal or cardiac muscle disorders, pulmonary disorders,  
 CC transplant rejection, disorders involving osteoclasts such as  
 CC osteoporosis, arthritis or malignancies, digestive/endocrine disorders,  
 CC infections and AIDS. The human secreted proteins of the invention are  
 CC represented in AAY07852-Y07993 and the encoding nucleic acids are  
 CC represented in AAX37451-X37552  
 XX

SQ Sequence 1434 BP; 480 A; 204 C; 250 G; 495 T; 0 U; 5 Other;

Alignment Scores:  
 Pred. No.: 7.13e-07 Length: 1434  
 Score: 98.00 Matches: 19  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 98.99% Indels: 0  
 DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AAX37452 (1-1434)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19  
 Db 507 ATGGCAGCCCAATTCAGCTTTGAGTTTCTTCTCTGGACACCTTATGCTCTGAAATCA 563

RESULT 2  
 ADA39771  
 ID ADA39771 standard; cDNA; 1434 BP.  
 XX  
 AC ADA39771;  
 XX  
 XX 20-NOV-2003 (first entry)  
 XX  
 XX Human secreted protein encoding cDNA.  
 XX  
 XX Human; secreted protein; cancer; hyperproliferative disorder;  
 KW rheumatoid arthritis; autoimmune disorder; haematopoietic disorder;  
 KW anaemia; allergic reaction; asthma; cardiovascular disorder;  
 KW wound healing; cytostatic; immunosuppressive; nootropic; neuroprotective;  
 KW antiviral; antiallergic; hepatotropic; antidiabetic; antiinflammatory;  
 KW vulnary; cardiant; gene therapy; ss.

OS Homo sapiens.  
 XX  
 PN WO2002102993-A2.  
 XX  
 XX 27-DEC-2002.  
 XX  
 XX 19-MAR-2002; 2002WO-US008123.  
 XX  
 XX 21-MAR-2001; 2001US-0277340P.  
 PR 19-JUL-2001; 2001US-0306171P.  
 PR 13-NOV-2001; 2001US-0331287P.  
 XX  
 XX (HUMA-) HUMAN GENOME SCI INC.  
 XX  
 XX Rosen CA, Ruben SM;  
 PI  
 XX WPI; 2003-175238/17.  
 XX  
 XX New human secreted proteins and nucleic acid molecules, useful for  
 PT preparing a diagnostic or pharmaceutical composition for diagnosing,  
 PT preventing or treating cancer or other hyperproliferative disorder,  
 PT asthma, allergies or AIDS.  
 XX

Claim 9; SEQ ID NO 153; 3205pp; English.

CC The invention relates to novel genes ADA39629-ADA40565 and proteins  
 CC ADA40566-ADA41501 for human secreted proteins, useful for preventing,  
 CC treating or ameliorating medical conditions e.g. by protein or gene  
 CC therapy. The polypeptides, nucleic acid molecules, antibodies or their  
 CC fragments, and agonists or antagonists that bind to the polypeptide are  
 CC useful for preparing a diagnostic or pharmaceutical composition for  
 CC diagnosing or treating cancer or other hyperproliferative disorder. The  
 CC polypeptides and nucleic acid molecules are also useful for detecting,  
 CC preventing, diagnosing, prognosticating, treating or ameliorating cancer  
 CC or other hyperproliferative disorders including neoplasms, autoimmune  
 CC disorders (e.g. diabetes, rheumatoid arthritis, systemic lupus  
 CC erythematosus, multiple sclerosis, autoimmune thyroiditis or haemolytic  
 CC anaemia), haematopoietic or haematological disorders (e.g. anaemia,  
 CC thrombocytopenia), allergic reactions including asthma or eczema,  
 CC inflammatory disorders (e.g. ischaemia-reperfusion injury, inflammatory  
 CC bowel disease or Crohn's disease), neurodegenerative disorders (e.g.  
 CC Alzheimer's disease or Parkinson's disease), cardiovascular disorders  
 CC (e.g. atherosclerosis, myocarditis), infectious diseases (bacterial,  
 CC fungal or viral infections including HIV/AIDS), or wound healing and  
 CC disorders of epithelial cell proliferation. The nucleic acids are also  
 CC useful for chromosome identification, radiation hybrid mapping or long-  
 CC range restriction mapping, as molecular weight markers, or as  
 CC hybridization or diagnostic probes. The polypeptides and antibodies are  
 CC useful for providing immunological probes for differential identification  
 CC of the tissues immunohistochemistry assays. Note: The sequence data for  
 CC this patent did not form part of the printed specification, but was  
 CC obtained in electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences.

XX  
 SQ Sequence 1434 BP; 480 A; 203 C; 250 G; 496 T; 0 U; 5 Other;

Alignment Scores:  
 Pred. No.: 7.13e-07 Length: 1434  
 Score: 98.00 Matches: 19  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 98.99% Indels: 0  
 DB: 7 Gaps: 0

US-10-799-747-116 (1-20) x ADA39771 (1-1434)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19  
 Db 507 ATGGCAGCCCAATTCAGCTTTGAGTTTCTTCTCTGGACACCTTATGCTCTGAAATCA 563  
 RESULT 3  
 ACC50424  
 ID ACC50424 standard; cDNA; 1434 BP.



PR 08-SEP-2000; 2000US-0232080P.  
PR 08-SEP-2000; 2000US-0232081P.  
PR 12-SEP-2000; 2000US-0231968P.  
PR 14-SEP-2000; 2000US-0232397P.  
PR 14-SEP-2000; 2000US-0232398P.  
PR 14-SEP-2000; 2000US-0232399P.  
PR 14-SEP-2000; 2000US-0232400P.  
PR 14-SEP-2000; 2000US-0232401P.  
PR 14-SEP-2000; 2000US-0233063P.  
PR 14-SEP-2000; 2000US-0233064P.  
PR 14-SEP-2000; 2000US-0233065P.  
PR 21-SEP-2000; 2000US-0234223P.  
PR 21-SEP-2000; 2000US-0234274P.  
PR 25-SEP-2000; 2000US-0234997P.  
PR 25-SEP-2000; 2000US-0234998P.  
PR 26-SEP-2000; 2000US-0235484P.  
PR 27-SEP-2000; 2000US-0235834P.  
PR 27-SEP-2000; 2000US-0235836P.  
PR 29-SEP-2000; 2000US-0236327P.  
PR 29-SEP-2000; 2000US-0236367P.  
PR 29-SEP-2000; 2000US-0236368P.  
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PR 29-SEP-2000; 2000US-0236370P.  
PR 02-OCT-2000; 2000US-0236802P.  
PR 02-OCT-2000; 2000US-0237037P.  
PR 02-OCT-2000; 2000US-0237038P.  
PR 02-OCT-2000; 2000US-0237039P.  
PR 02-OCT-2000; 2000US-0237040P.  
PR 13-OCT-2000; 2000US-0239335P.  
PR 13-OCT-2000; 2000US-0239337P.  
PR 20-OCT-2000; 2000US-0240960P.  
PR 20-OCT-2000; 2000US-0241221P.  
PR 20-OCT-2000; 2000US-0241785P.  
PR 20-OCT-2000; 2000US-0241786P.  
PR 20-OCT-2000; 2000US-0241787P.  
PR 20-OCT-2000; 2000US-0241808P.  
PR 20-OCT-2000; 2000US-0241809P.  
PR 20-OCT-2000; 2000US-0241826P.  
PR 01-NOV-2000; 2000US-0244617P.  
PR 08-NOV-2000; 2000US-0246474P.  
PR 08-NOV-2000; 2000US-0246475P.  
PR 08-NOV-2000; 2000US-0246476P.  
PR 08-NOV-2000; 2000US-0246477P.  
PR 08-NOV-2000; 2000US-0246478P.  
PR 08-NOV-2000; 2000US-0246523P.  
PR 08-NOV-2000; 2000US-0246524P.  
PR 08-NOV-2000; 2000US-0246525P.  
PR 08-NOV-2000; 2000US-0246526P.  
PR 08-NOV-2000; 2000US-0246527P.  
PR 08-NOV-2000; 2000US-0246528P.  
PR 08-NOV-2000; 2000US-0246532P.  
PR 08-NOV-2000; 2000US-0246609P.  
PR 08-NOV-2000; 2000US-0246610P.  
PR 08-NOV-2000; 2000US-0246611P.  
PR 08-NOV-2000; 2000US-0246613P.  
PR 17-NOV-2000; 2000US-0249207P.  
PR 17-NOV-2000; 2000US-0249208P.  
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PR 17-NOV-2000; 2000US-0249210P.  
PR 17-NOV-2000; 2000US-0249211P.  
PR 17-NOV-2000; 2000US-0249212P.  
PR 17-NOV-2000; 2000US-0249213P.  
PR 17-NOV-2000; 2000US-0249214P.  
PR 17-NOV-2000; 2000US-0249215P.  
PR 17-NOV-2000; 2000US-0249216P.  
PR 17-NOV-2000; 2000US-0249217P.  
PR 17-NOV-2000; 2000US-0249218P.  
PR 17-NOV-2000; 2000US-0249218P.  
PR 17-NOV-2000; 2000US-0249244P.  
PR 17-NOV-2000; 2000US-0249245P.  
PR 17-NOV-2000; 2000US-0249264P.  
PR 17-NOV-2000; 2000US-0249265P.  
PR 17-NOV-2000; 2000US-0249297P.  
PR 17-NOV-2000; 2000US-0249299P.

PR 17-NOV-2000; 2000US-0249300P.  
PR 01-DEC-2000; 2000US-0250160P.  
PR 01-DEC-2000; 2000US-0250391P.  
PR 05-DEC-2000; 2000US-0251030P.  
PR 05-DEC-2000; 2000US-0251988P.  
PR 05-DEC-2000; 2000US-0256719P.  
PR 06-DEC-2000; 2000US-0251479P.  
PR 08-DEC-2000; 2000US-0251856P.  
PR 08-DEC-2000; 2000US-0251868P.  
PR 08-DEC-2000; 2000US-0251869P.  
PR 08-DEC-2000; 2000US-0251989P.  
PR 08-DEC-2000; 2000US-0251990P.  
PR 11-DEC-2000; 2000US-0254097P.  
PR 05-JAN-2001; 2001US-0259678P.  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX Rosen CA, Barash SC, Ruben SM;  
XX WPI; 2001-502630/55.  
XX Polynucleotides encoding digestive system antigens, useful for  
PT diagnosing, treating, preventing and/or prognosing disorders of the  
PT digestive system, particularly cancer and cancer metastases.  
XX Disclosure; SEQ ID NO 3071; 986pp; English.  
XX The present invention provides the protein and coding sequences of a  
CC number of human digestive system antigens. These can be used in the  
CC diagnosis, treatment and prevention of digestive system disorders,  
CC including cancer, Meckel's diverticulum, bacterial or parasitic  
CC infections, appendicitis, Hirschsprung's disease, chronic colitis or  
CC ulcerative colitis. The present sequence is a genomic DNA fragment  
CC encoding a digestive system antigen of the invention  
XX  
SQ Sequence 8417 BP; 2708 A; 1429 C; 1657 G; 2621 T; 0 U; 2 Other;  
  
Alignment Scores:  
Pred. No.: 102 Length: 8417  
Score: 56.00 Matches: 9  
Percent Similarity: 78.57% Conservative: 2  
Best Local Similarity: 64.29% Mismatches: 3  
Query Match: 56.57% Indels: 0  
DB: 4 Gaps: 0  
  
US-10-799-747-116 (1-20) x AAK89495 (1-8417)  
  
QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTyrThrPro 14  
||| |||||:||||:||||| |||||  
Db 7484 ATGCCACAACTTCGATTTGACGTTTATTCTGCACCCA 7443  
  
RESULT 5  
AAS32681  
ID AAS32681 standard; DNA; 8417 BP.  
XX  
AC AAS32681;  
XX  
DT 17-DEC-2001 (first entry)  
XX  
DE Human genomic DNA for novel endocrine antigen, SEQ ID No 635.  
XX  
KW Human; endocrine antigen; ds; cytostatic; antiinfertility; antidiabetic;  
KW thyroid-active; adrenal-active; androgenic; gastric; gene therapy;  
KW antitense-therapy; antibody; endocrine disorder; hormone imbalance;  
KW reproductive disorder; endocrine cancer; pancreatic disorder;  
KW diabetes mellitus; adrenal gland disorder; hirsutism; thyroid disorder;  
KW hyperthyroidism; hypothalamic disorder; vanishing testes syndrome.  
XX Homo sapiens.  
XX  
XX WC200155319-A2.  
XX  
PD 02-AUG-2001.



XX 17-JAN-2001; 2001WO-US0001335.  
PF 31-JAN-2000; 2000US-0179065P.  
XX 04-FEB-2000; 2000US-0180628P.  
PR 24-FEB-2000; 2000US-0184664P.  
PR 02-MAR-2000; 2000US-0186350P.  
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PR 17-MAR-2000; 2000US-0190076P.  
PR 18-APR-2000; 2000US-0198123P.  
PR 19-MAY-2000; 2000US-0205151P.  
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PR 28-JUN-2000; 2000US-0214886P.  
PR 30-JUN-2000; 2000US-0215135P.  
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PR 08-SEP-2000; 2000US-0232080P.  
PR 08-SEP-2000; 2000US-0232081P.  
PR 12-SEP-2000; 2000US-0231968P.  
PR 14-SEP-2000; 2000US-0232397P.  
PR 14-SEP-2000; 2000US-0232398P.  
PR 14-SEP-2000; 2000US-0232399P.  
PR 14-SEP-2000; 2000US-0232400P.  
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PR 14-SEP-2000; 2000US-0233063P.  
PR 14-SEP-2000; 2000US-0233064P.  
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PR 21-SEP-2000; 2000US-0234274P.  
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PR 29-SEP-2000; 2000US-0236367P.  
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PR 20-OCT-2000; 2000US-0241786P.  
PR 20-OCT-2000; 2000US-0241787P.  
PR 20-OCT-2000; 2000US-0241808P.  
PR 20-OCT-2000; 2000US-0241809P.  
PR 20-OCT-2000; 2000US-0241826P.  
PR 01-NOV-2000; 2000US-0244617P.  
PR 08-NOV-2000; 2000US-0246474P.  
PR 08-NOV-2000; 2000US-0246475P.  
PR 08-NOV-2000; 2000US-0246476P.  
PR 08-NOV-2000; 2000US-0246477P.  
PR 08-NOV-2000; 2000US-0246478P.  
PR 08-NOV-2000; 2000US-0246523P.  
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PR 08-NOV-2000; 2000US-0246528P.  
PR 08-NOV-2000; 2000US-0246532P.  
PR 08-NOV-2000; 2000US-0246609P.  
PR 08-NOV-2000; 2000US-0246610P.  
PR 08-NOV-2000; 2000US-0246611P.  
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PR 17-NOV-2000; 2000US-0249207P.  
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PR 17-NOV-2000; 2000US-0249244P.  
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PR 17-NOV-2000; 2000US-0249246P.  
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PR 01-DEC-2000; 2000US-0249300P.  
PR 01-DEC-2000; 2000US-0250160P.  
PR 01-DEC-2000; 2000US-0250391P.  
PR 05-DEC-2000; 2000US-0251030P.  
PR 05-DEC-2000; 2000US-0251988P.  
PR 06-DEC-2000; 2000US-0256719P.  
PR 08-DEC-2000; 2000US-0251479P.  
PR 08-DEC-2000; 2000US-0251856P.  
PR 08-DEC-2000; 2000US-0251868P.  
PR 08-DEC-2000; 2000US-0251869P.  
PR 08-DEC-2000; 2000US-0251989P.  
PR 11-DEC-2000; 2000US-0251990P.  
PR 05-JAN-2001; 2000US-0254097P.  
PR 05-JAN-2001; 2000US-0259678P.  
(HUMA-) HUMAN GENOME SCI INC.  
PA Rosen CA, Barash SC, Ruben SM;  
PI WPI; 2001-457726/49.  
XX  
XX  
XX  
XX  
XX

PT Isolated polypeptide for treating, preventing and prognosing disorders  
 PT related to the endocrine system including endocrine disorders,  
 PT reproductive disorders, and gastrointestinal disorders and also for  
 PT testing and detection e.g. diagnosis.

XX PS Disclosure; SEQ ID NO 635; 558pp; English.

XX CC The invention relates to cDNAs encoding novel human endocrine antigens or  
 CC a fragment having biological activity, a domain, an epitope, full length  
 CC protein, variant, allelic variant or a species homologue of the  
 CC cDNA/antigen. The DNAs and polypeptides are useful for preventing,  
 CC treating or ameliorating a medical condition when administered (e.g. by  
 CC gene therapy or antisense-therapy). Identifying mutations in the genes  
 CC coding for the antigens is useful for diagnosing a pathological condition  
 CC or a susceptibility to a pathological condition. The DNAs, antigens and  
 CC antibodies raised against the antigens useful for treating, preventing  
 CC and/or prognosing disorders related to the endocrine system or hormone  
 CC imbalance or reproductive disorders, cancers of endocrine tissues,  
 CC disorders of the pancreas (e.g. diabetes mellitus), the adrenal glands  
 CC (e.g. hirsutism), ovaries, the thyroid (e.g. hyperthyroidism), the  
 CC hypothalamus and testes (e.g. vanishing testes syndrome), many examples  
 CC of diseases and disorders are given in the specification. The present  
 CC sequence is genomic DNA fragment form a gene encoding an endocrine  
 CC antigen of the invention. Note: The sequence data for this patent did not  
 CC form part of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences

XX SQ Sequence 8417 BP; 2621 A; 1657 C; 1429 G; 2708 T; 0 U; 2 Other;

Alignment Scores:  
 Pred. No.: 102 Length: 8417  
 Score: 56.00 Matches: 9  
 Percent Similarity: 78.57% Conservative: 2  
 Best Local Similarity: 64.29% Mismatches: 3  
 Query Match: 56.57% Indels: 0  
 DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x AAS32681 (1-8417)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTrpThrPro 14

DB 934 ATGCCACACATTCGATTGACGTTTATTCTCGACCCCA 975

RESULT 6

AAS32472

ID AAS32472 standard; DNA; 5760 BP.

XX AC AAS32472;

XX DT 17-DEC-2001 (first entry)

XX DE Human genomic DNA for novel endocrine antigen, SEQ ID No 426.

XX KW Human; endocrine antigen; ds; cytostatic; antiinfertility; antidiabetic;  
 KW thyroid-active; adrenal-active; androgenic; gastric; gene therapy;  
 KW antisense-therapy; antibody; endocrine disorder; hormone imbalance;  
 KW reproductive disorder; endocrine cancer; pancreatic disorder;  
 KW diabetes mellitus; adrenal gland disorder; hirsutism; thyroid disorder;  
 KW hyperthyroidism; hypothalamic disorder; vanishing testes syndrome.

XX OS Homo sapiens.

XX PN WC200155319-A2.

XX PD 02-AUG-2001.

XX PF 17-JAN-2001; 2001WO-US0001335.

XX PR 31-JAN-2000; 2000US-0179065P.

PR 04-FEB-2000; 2000US-0180628P.

PR 24-FEB-2000; 2000US-0184664P.

PR 02-MAR-2000; 2000US-0186350P.

PR 16-MAR-2000; 2000US-0189874P.

PR 17-MAR-2000; 2000US-0190076P.  
 PR 18-APR-2000; 2000US-0198123P.  
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 PR 28-JUN-2000; 2000US-0214886P.  
 PR 30-JUN-2000; 2000US-0215135P.  
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 PR 11-JUL-2000; 2000US-0217487P.  
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 PR 18-AUG-2000; 2000US-0226279P.  
 PR 22-AUG-2000; 2000US-0226681P.  
 PR 22-AUG-2000; 2000US-0226868P.  
 PR 22-AUG-2000; 2000US-0227182P.  
 PR 23-AUG-2000; 2000US-0227009P.  
 PR 30-AUG-2000; 2000US-0228924P.  
 PR 01-SEP-2000; 2000US-0229287P.  
 PR 01-SEP-2000; 2000US-0229343P.  
 PR 01-SEP-2000; 2000US-0229344P.  
 PR 01-SEP-2000; 2000US-0229345P.  
 PR 05-SEP-2000; 2000US-0229509P.  
 PR 05-SEP-2000; 2000US-0229513P.  
 PR 06-SEP-2000; 2000US-0230437P.  
 PR 06-SEP-2000; 2000US-0230438P.  
 PR 08-SEP-2000; 2000US-0231242P.  
 PR 08-SEP-2000; 2000US-0231243P.  
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 PR 08-SEP-2000; 2000US-0231413P.  
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 PR 08-SEP-2000; 2000US-0232080P.  
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 PR 12-SEP-2000; 2000US-0231968P.  
 PR 14-SEP-2000; 2000US-0232397P.  
 PR 14-SEP-2000; 2000US-0232398P.  
 PR 14-SEP-2000; 2000US-0232399P.  
 PR 14-SEP-2000; 2000US-0232400P.  
 PR 14-SEP-2000; 2000US-0232401P.  
 PR 14-SEP-2000; 2000US-0233063P.  
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 PR 21-SEP-2000; 2000US-0234223P.  
 PR 21-SEP-2000; 2000US-0234274P.  
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 PR 26-SEP-2000; 2000US-0235484P.  
 PR 27-SEP-2000; 2000US-0235834P.  
 PR 27-SEP-2000; 2000US-0235836P.  
 PR 29-SEP-2000; 2000US-0236327P.  
 PR 29-SEP-2000; 2000US-0236367P.  
 PR 29-SEP-2000; 2000US-0236368P.  
 PR 29-SEP-2000; 2000US-0236369P.  
 PR 29-SEP-2000; 2000US-0236370P.  
 PR 02-OCT-2000; 2000US-0236802P.  
 PR 02-OCT-2000; 2000US-0237037P.  
 PR 02-OCT-2000; 2000US-0237038P.  
 PR 02-OCT-2000; 2000US-0237039P.  
 PR 02-OCT-2000; 2000US-0237040P.  
 PR 13-OCT-2000; 2000US-0239935P.

PR 13-OCT-2000; 2000US-0239937P.  
PR 20-OCT-2000; 2000US-0240960P.  
PR 20-OCT-2000; 2000US-0241221P.  
PR 20-OCT-2000; 2000US-0241785P.  
PR 20-OCT-2000; 2000US-0241786P.  
PR 20-OCT-2000; 2000US-0241787P.  
PR 20-OCT-2000; 2000US-0241808P.  
PR 20-OCT-2000; 2000US-0241809P.  
PR 20-OCT-2000; 2000US-0241826P.  
PR 01-NOV-2000; 2000US-0244617P.  
PR 08-NOV-2000; 2000US-0246474P.  
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PR 08-NOV-2000; 2000US-0246478P.  
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PR 08-NOV-2000; 2000US-0246528P.  
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PR 08-NOV-2000; 2000US-0246609P.  
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PR 08-NOV-2000; 2000US-0246611P.  
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PR 17-NOV-2000; 2000US-0249207P.  
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PR 17-NOV-2000; 2000US-0249300P.  
PR 01-DEC-2000; 2000US-0250160P.  
PR 01-DEC-2000; 2000US-0250391P.  
PR 05-DEC-2000; 2000US-0251030P.  
PR 05-DEC-2000; 2000US-0251988P.  
PR 05-DEC-2000; 2000US-0256719P.  
PR 06-DEC-2000; 2000US-0251479P.  
PR 06-DEC-2000; 2000US-0251856P.  
PR 08-DEC-2000; 2000US-0251868P.  
PR 08-DEC-2000; 2000US-0251869P.  
PR 08-DEC-2000; 2000US-0251989P.  
PR 08-DEC-2000; 2000US-0251990P.  
PR 11-DEC-2000; 2000US-0254097P.  
PR 05-JAN-2001; 2001US-0259678P.  
XX (HUMA-) HUMAN GENOME SCI INC.  
XX  
XX Rosen CA, Barash SC, Ruben SM;  
XX WPI; 2001-457726/49.  
XX  
XX Isolated polypeptide for treating, preventing and prognosing disorders  
XX related to the endocrine system including endocrine disorders,  
XX reproductive disorders, and gastrointestinal disorders and also for  
XX testing and detection e.g. diagnosis.  
XX  
XX Disclosure; SEQ ID NO 426; 558pp; English.  
XX  
XX The invention relates to cDNAs encoding novel human endocrine antigens or

CC a fragment having biological activity, a domain, an epitope, full length  
CC protein, variant, allelic variant or a species homologue of the  
CC cDNA/antigen. The DNAs and polypeptides are useful for preventing,  
CC treating or ameliorating a medical condition when administered (e.g. by  
CC gene therapy or antisense-therapy). Identifying mutations in the genes  
CC coding for the antigens is useful for diagnosing a pathological condition  
CC or a susceptibility to a pathological condition. The DNAs, antigens and  
CC antibodies raised against the antigens useful for treating, preventing  
CC and/or prognosing disorders related to the endocrine system or hormone  
CC imbalance or reproductive disorders, cancers of endocrine tissues,  
CC disorders of the pancreas (e.g. diabetes mellitus), the adrenal glands  
CC (e.g. hirsutism), ovaries, the thyroid (e.g. hyperthyroidism), the  
CC hypothalamus and testes (e.g. vanishing testes syndrome), many examples  
CC of diseases and disorders are given in the specification. The present  
CC sequence is genomic DNA fragment form a gene encoding an endocrine  
CC antigen of the invention. Note: The sequence data for this patent did not  
CC form part of the printed specification, but was obtained in electronic  
CC format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
XX  
SQ Sequence 5760 BP; 1725 A; 1199 C; 1068 G; 1768 T; 0 U; 0 Other;  
  
Alignment Scores:  
Pred. No.: 208 Length: 5760  
Score: 53.00 Matches: 10  
Percent Similarity: 68.75% Conservative: 1  
Best Local Similarity: 62.50% Mismatches: 5  
Query Match: 53.54% Indels: 0  
DB: 4 Gaps: 0  
  
US-10-799-747-116 (1-20) x AAS32472 (1-5760)  
  
QY 2 AlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeu 17  
Db 567 GCCCGGCACCTCAACCTTGAGCTTTGGCTTGGGTGTATATCTT 614  
  
RESULT 7  
AAS32471  
ID AAS32471 standard; DNA; 5763 BP.  
XX  
AC AAS32471;  
XX  
DT 17-DEC-2001 (first entry)  
XX  
DE Human genomic DNA for novel endocrine antigen, SEQ ID No 425.  
XX  
KW Human; endocrine antigen; ds; cytostatic; antiinfertility; antidiabetic;  
KW thyroid-active; adrenal-active; androgenic; gastric; gene therapy;  
KW antisense-therapy; antibody; endocrine disorder; hormone imbalance;  
KW reproductive disorder; endocrine cancer; pancreatic disorder;  
KW diabetes mellitus; adrenal gland disorder; hirsutism; thyroid disorder;  
KW hyperthyroidism; hypothalamic disorder; vanishing testes syndrome.  
XX  
OS Homo sapiens.  
XX  
PN WO200155319-A2.  
XX  
PD 02-AUG-2001.  
XX  
PF 17-JAN-2001; 2001WO-US001335.  
XX  
PR 31-JAN-2000; 2000US-0179065P.  
PR 04-FEB-2000; 2000US-0180628P.  
PR 24-FEB-2000; 2000US-0184664P.  
PR 02-MAR-2000; 2000US-0186350P.  
PR 16-MAR-2000; 2000US-0189874P.  
PR 17-MAR-2000; 2000US-0190076P.  
PR 18-APR-2000; 2000US-0198123P.  
PR 19-MAY-2000; 2000US-0205515P.  
PR 07-JUN-2000; 2000US-0209467P.  
PR 28-JUN-2000; 2000US-0214886P.  
PR 30-JUN-2000; 2000US-0215135P.  
PR 07-JUL-2000; 2000US-0216647P.  
PR 07-JUL-2000; 2000US-0216880P.



CC and/ or prognosing disorders related to the endocrine system or hormone  
 CC imbalance or reproductive disorders, cancers of endocrine tissues,  
 CC disorders of the pancreas (e.g. diabetes mellitus), the adrenal glands  
 CC (e.g. hirsutism), ovaries, the thyroid (e.g. hyperthyroidism), the  
 CC hypothalamus and testes (e.g. vanishing testes syndrome), many examples  
 CC of diseases and disorders are given in the specification. The present  
 CC sequence is genomic DNA fragment form a gene encoding an endocrine  
 CC antigen of the invention. Note: The sequence data for this patent did not  
 CC form part of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 SQ Sequence 5763 BP; 1725 A; 1203 C; 1066 G; 1768 T; 0 U; 1 Other;

Alignment Scores:  
 Pred. No.: 209 Length: 5763  
 Score: 53.00 Matches: 10  
 Percent Similarity: 68.75% Conservative: 1  
 Best Local Similarity: 62.50% Mismatches: 5  
 Query Match: 53.54% Indels: 0  
 DB: 4 Gaps: 0  
 US-10-799-747-116 (1-20) x AAS32471 (1-5763)

QY 2 AlaAlaHisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17  
 DB 567 GCCCGCAGCTCAACCTGAGCTTTGCCCTTGGTGTATATCTCTT 614

RESULT 8  
 ABZ36220  
 ID ABZ36220 standard; cDNA; 2877 BP.  
 AC ABZ36220;  
 XX  
 DT 10-FEB-2003 (first entry)  
 XX  
 DE Human secretory polynucleotide SPTM SEQ ID NO 384.  
 XX  
 KW Human; SPTM; autoimmune disorder; inflammatory disorder; AIDS; anaemia;  
 KW asthma; Crohn's disease; neurological disorder; epilepsy; cancer;  
 KW Huntington's disease; Alzheimer's disease; Creutzfeldt-Jakob disease;  
 KW multiple sclerosis; Parkinson's disease; cell proliferative disorder;  
 KW anti-inflammatory; immunosuppressive; neuroprotective; nootropic;  
 KW neuroleptic; anticonvulsant; cytostatic; antiparkinsonian; anxiolytic;  
 KW antipsoriatic; antianaemic; anti-HIV; human immunodeficiency virus;  
 KW secretory polynucleotide; secretory protein; gene; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200283876-A2.  
 XX  
 PD 24-OCT-2002.  
 XX  
 PF 27-MAR-2002; 2002WO-US0009921.  
 XX  
 PR 29-MAR-2001; 2001US-0280067P.  
 PR 29-MAR-2001; 2001US-0280068P.  
 PR 16-MAY-2001; 2001US-0291280P.  
 PR 17-MAY-2001; 2001US-0291829P.  
 PR 17-MAY-2001; 2001US-0291849P.  
 PR 19-JUN-2001; 2001US-0299428P.  
 PR 20-JUN-2001; 2001US-0299776P.  
 PR 20-JUN-2001; 2001US-030001P.  
 XX  
 PA (INCY-) INCYTE GENOMICS INC.  
 XX  
 PI Daffo A, Jones AL, Tran AB, Dahl CR, Gietzen D, Chinn J;  
 PI Dufour GE, Hillman JL, Yu JY, Thason O, Yap PE, Anshey SR;  
 PI Daugherty SC, Dam TC, Liu IF, Nguyen DA, Kleefeld Y, Gerstin EH;  
 PI Peralta CH, David ME, Lewis SA, Chen AJ, Panzer SR, Harris B;  
 PI Flores V, Marwaha R, Lo A, Lan RY, Urashka ME;  
 XX  
 DR WPI; 2003-075543/07.  
 DR P-PSDB; ABP75778.

XX New human secretory proteins and polynucleotides, useful for diagnosing,  
 PT treating or preventing autoimmune/inflammatory disorders (e.g. AIDS),  
 PT neurological disorders (e.g. Alzheimer's), or cell proliferations or  
 PT cancers.  
 XX  
 PS Claim 1; SEQ ID NO 384; 458pp + Sequence Listing; English.  
 XX  
 CC The invention relates to a secretory polynucleotide (designated sptm)  
 CC comprising any of 567 polynucleotide sequences (ABZ35837-ABZ36403), a  
 CC naturally occurring polynucleotide sequence at least 90 % identical to  
 CC the polynucleotide sequence, a polynucleotide complementary to them or an  
 CC RNA equivalent of them. The polypeptide or polynucleotide are useful for  
 CC treating, preventing or diagnosing a disease or condition associated with  
 CC the expression of functional SPTM. These are particularly useful for  
 CC diagnosing, treating or preventing autoimmune/inflammatory disorders  
 CC (e.g. acquired immunodeficiency syndrome, anaemia, asthma or Crohn's  
 CC disease), neurological disorders (e.g. epilepsy, Huntington's disease,  
 CC dementia, stroke, Alzheimer's disease, Creutzfeldt-Jakob disease,  
 CC multiple sclerosis, cerebral palsy, Parkinson's disease, anxiety,  
 CC schizophrenia or amnesia), or cell proliferative disorders (e.g.  
 CC psoriasis, polycythemia vera, or cancers including adenocarcinoma,  
 CC leukaemia, lymphoma, melanoma, myeloma, sarcoma or cancers of the brain,  
 CC breast, cervix or prostate). Note: The sequence data for this patent did  
 CC not form part of the printed specification, but was obtained in electronic  
 CC format directly from WIPO at ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 SQ Sequence 2877 BP; 854 A; 609 C; 504 G; 910 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 131 Length: 2877  
 Score: 52.00 Matches: 9  
 Percent Similarity: 73.33% Conservative: 2  
 Best Local Similarity: 60.00% Mismatches: 4  
 Query Match: 52.53% Indels: 0  
 DB: 7 Gaps: 0

US-10-799-747-116 (1-20) x ABZ36220 (1-2877)

QY 2 AlaAlaHisSerValLeuSerPheLeuLeuTTPThrProTyrAla 16  
 DB 2193 GCTCTCAGCAAAATATCATTCATTATTTGGTGCCACAGGCC 2237

RESULT 9  
 AAK82208/c  
 ID AAK82208 standard; DNA; 36785 BP.  
 AC AAK82208;  
 XX  
 DT 07-NOV-2001 (first entry)  
 XX  
 DE Human immune/haematopoietic antigen genomic sequence SEQ ID NO:37020.  
 XX  
 KW Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
 KW cytostatic; gene therapy; vaccine; metastasis; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200157182-A2.  
 XX  
 PD 09-AUG-2001.  
 XX  
 PF 17-JAN-2001; 2001WO-US001354.  
 XX  
 PR 31-JAN-2000; 2000US-0179065P.  
 PR 04-FEB-2000; 2000US-0180628P.  
 PR 24-FEB-2000; 2000US-0184664P.  
 PR 02-MAR-2000; 2000US-0186350P.  
 PR 16-MAR-2000; 2000US-0189874P.  
 PR 17-MAR-2000; 2000US-0190076P.  
 PR 18-APR-2000; 2000US-0198123P.  
 PR 19-MAY-2000; 2000US-0205515P.  
 PR 07-JUN-2000; 2000US-0209467P.

PR 28-JUN-2000; 2000US-0214886P.  
PR 30-JUN-2000; 2000US-0215135P.  
PR 07-JUL-2000; 2000US-0216647P.  
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PR 11-JUL-2000; 2000US-0217487P.  
PR 11-JUL-2000; 2000US-0217496P.  
PR 14-JUL-2000; 2000US-0218290P.  
PR 26-JUL-2000; 2000US-0220963P.  
PR 26-JUL-2000; 2000US-0220964P.  
PR 14-AUG-2000; 2000US-0224518P.  
PR 14-AUG-2000; 2000US-0224519P.  
PR 14-AUG-2000; 2000US-0225213P.  
PR 14-AUG-2000; 2000US-0225214P.  
PR 14-AUG-2000; 2000US-0225266P.  
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PR 14-AUG-2000; 2000US-0225268P.  
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PR 22-AUG-2000; 2000US-0227182P.  
PR 23-AUG-2000; 2000US-0227009P.  
PR 30-AUG-2000; 2000US-0228524P.  
PR 01-SEP-2000; 2000US-0229287P.  
PR 01-SEP-2000; 2000US-0229343P.  
PR 01-SEP-2000; 2000US-0229344P.  
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PR 05-SEP-2000; 2000US-0229509P.  
PR 05-SEP-2000; 2000US-0229513P.  
PR 06-SEP-2000; 2000US-0230437P.  
PR 06-SEP-2000; 2000US-0230438P.  
PR 08-SEP-2000; 2000US-02311242P.  
PR 08-SEP-2000; 2000US-02311243P.  
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PR 08-SEP-2000; 2000US-0231414P.  
PR 08-SEP-2000; 2000US-0232080P.  
PR 08-SEP-2000; 2000US-0232081P.  
PR 12-SEP-2000; 2000US-0231968P.  
PR 14-SEP-2000; 2000US-0232397P.  
PR 14-SEP-2000; 2000US-0232398P.  
PR 14-SEP-2000; 2000US-0232399P.  
PR 14-SEP-2000; 2000US-0232400P.  
PR 14-SEP-2000; 2000US-0232401P.  
PR 14-SEP-2000; 2000US-0233063P.  
PR 14-SEP-2000; 2000US-0233064P.  
PR 14-SEP-2000; 2000US-0233065P.  
PR 21-SEP-2000; 2000US-0234223P.  
PR 21-SEP-2000; 2000US-0234274P.  
PR 25-SEP-2000; 2000US-0234997P.  
PR 25-SEP-2000; 2000US-0234998P.  
PR 26-SEP-2000; 2000US-0235484P.  
PR 27-SEP-2000; 2000US-0235834P.  
PR 27-SEP-2000; 2000US-0235836P.  
PR 29-SEP-2000; 2000US-0236327P.  
PR 29-SEP-2000; 2000US-0236367P.  
PR 29-SEP-2000; 2000US-0236368P.  
PR 29-SEP-2000; 2000US-0236369P.  
PR 29-SEP-2000; 2000US-0236370P.  
PR 02-OCT-2000; 2000US-0236802P.  
PR 02-OCT-2000; 2000US-0237037P.  
PR 02-OCT-2000; 2000US-0237038P.  
PR 02-OCT-2000; 2000US-0237039P.  
PR 02-OCT-2000; 2000US-0237040P.  
PR 13-OCT-2000; 2000US-0239935P.  
PR 13-OCT-2000; 2000US-0239937P.  
PR 20-OCT-2000; 2000US-0240960P.  
PR 20-OCT-2000; 2000US-0241221P.  
PR 20-OCT-2000; 2000US-0241785P.

PR 20-OCT-2000; 2000US-0241786P.  
PR 20-OCT-2000; 2000US-0241787P.  
PR 20-OCT-2000; 2000US-0241808P.  
PR 20-OCT-2000; 2000US-0241809P.  
PR 20-OCT-2000; 2000US-0241826P.  
PR 01-NOV-2000; 2000US-0244617P.  
PR 08-NOV-2000; 2000US-0246474P.  
PR 08-NOV-2000; 2000US-0246475P.  
PR 08-NOV-2000; 2000US-0246476P.  
PR 08-NOV-2000; 2000US-0246477P.  
PR 08-NOV-2000; 2000US-0246478P.  
PR 08-NOV-2000; 2000US-0246523P.  
PR 08-NOV-2000; 2000US-0246524P.  
PR 08-NOV-2000; 2000US-0246525P.  
PR 08-NOV-2000; 2000US-0246526P.  
PR 08-NOV-2000; 2000US-0246527P.  
PR 08-NOV-2000; 2000US-0246528P.  
PR 08-NOV-2000; 2000US-0246532P.  
PR 08-NOV-2000; 2000US-0246609P.  
PR 08-NOV-2000; 2000US-0246610P.  
PR 08-NOV-2000; 2000US-0246611P.  
PR 08-NOV-2000; 2000US-0246613P.  
PR 17-NOV-2000; 2000US-0249207P.  
PR 17-NOV-2000; 2000US-0249208P.  
PR 17-NOV-2000; 2000US-0249209P.  
PR 17-NOV-2000; 2000US-0249210P.  
PR 17-NOV-2000; 2000US-0249211P.  
PR 17-NOV-2000; 2000US-0249212P.  
PR 17-NOV-2000; 2000US-0249213P.  
PR 17-NOV-2000; 2000US-0249214P.  
PR 17-NOV-2000; 2000US-0249215P.  
PR 17-NOV-2000; 2000US-0249216P.  
PR 17-NOV-2000; 2000US-0249217P.  
PR 17-NOV-2000; 2000US-0249218P.  
PR 17-NOV-2000; 2000US-0249244P.  
PR 17-NOV-2000; 2000US-0249245P.  
PR 17-NOV-2000; 2000US-0249264P.  
PR 17-NOV-2000; 2000US-0249265P.  
PR 17-NOV-2000; 2000US-0249297P.  
PR 17-NOV-2000; 2000US-0249299P.  
PR 17-NOV-2000; 2000US-0249300P.  
PR 01-DEC-2000; 2000US-0250160P.  
PR 01-DEC-2000; 2000US-0250391P.  
PR 05-DEC-2000; 2000US-0251030P.  
PR 05-DEC-2000; 2000US-0251988P.  
PR 05-DEC-2000; 2000US-0256719P.  
PR 06-DEC-2000; 2000US-0251479P.  
PR 08-DEC-2000; 2000US-0251856P.  
PR 08-DEC-2000; 2000US-0251868P.  
PR 08-DEC-2000; 2000US-0251869P.  
PR 08-DEC-2000; 2000US-0251989P.  
PR 08-DEC-2000; 2000US-0251990P.  
PR 11-DEC-2000; 2000US-0254097P.  
PR 05-JAN-2001; 2001US-0259678P.

(HUMA-) HUMAN GENOME SCI INC.

Rosen CA, Barash SC, Ruben SM;

WPI; 2001-483426/52.

Nucleic acids encoding human immune/hematopoietic antigen polypeptides, useful for preventing, diagnosing and/or treating cancers and metastasis.

Disclosure; SEQ ID NO 37020; 3071pp + Sequence Listing; English.

AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I) amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic activity, and can be used in gene therapy and vaccine production. (I) proteins and polynucleotides may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate (I) expression. For example, they may be used to treat disorders associated with decreased expression by rectifying mutations or deletions in a patient's genome

CC that affect the activity of (I) by expressing inactive proteins or to  
 CC supplement the patients own production of (I). Additionally, (I)  
 CC polynucleotides may be used to produce the secreted (I), by inserting the  
 CC nucleic acids into a host cell and culturing the cell to express the  
 CC protein. (I) proteins and polynucleotides may be used to prevent,  
 CC diagnose and treat immune/haematopoietic-related diseases, especially  
 CC cancers and cancer metastases of haematopoietic-derived cells. AAK64703  
 CC to AAK87694 represent human immune/haematopoietic antigen genomic  
 CC sequences from the present invention. AAK54942 to AAK54950 and AAK82169  
 CC represent sequences used in the exemplification of the present invention  
 XX

SQ Sequence 36785 BP; 9938 A; 8156 C; 8241 G; 10450 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 3.04e+03 Length: 36785  
 Score: 52.00 Matches: 9  
 Percent Similarity: 73.33% Conservative: 2  
 Best Local Similarity: 60.00% Mismatches: 4  
 Query Match: 52.53% Indels: 0  
 DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x AAK82208 (1-36785)

QY 2 AlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAla 16

Db 13072 GCTCCTCAGACGAAATATCATTCATATTTGGTGCCACAGGCC 13028

RESULT 10

AAH64369/c

ID AAF64369 standard; cDNA; 400 BP.

XX AAF64369;

XX

XX

DT 09-APR-2001 (first entry)

XX

DE Novel human polynucleotide, SEQ ID NO: 125.

XX

KW Human; cytostatic; gene therapy; colon cancer; prostate cancer;  
 breast cancer; lung cancer; cancer detection; ss.

XX

OS Homo sapiens.

XX

PN WO200102568-A2.

XX

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PD 11-JAN-2001.

XX

PF 30-JUN-2000; 2000MO-US018374.

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PR 02-JUL-1999; 99US-0142310P.

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PR 02-JUL-1999; 99US-0142311P.

XX

PA (CHTR ) CHIRON CORP.

XX

PA (HYSE-) HYSEQ INC.

XX

PI Williams LT, Escobedo J, Inniss MA, Garcia PD, Klinger J;

XX

PI Kassam A, Reinhard C, Randazzo E, Kennedy GC, Pot D, Lamson G;

XX

PI Drmanac R, C-kenjakov R, Drmanac S, Dickson M, Labat I;

XX

PI Leshkowitz D, Kita D, Garcia V, Jones LW, Strache-Crain B;

XX

DR WPI; 2001-091805/10.

XX

PT Library of polynucleotides for diagnosing a cancerous state of a

XX

PT mammalian cell and detecting cancer, particularly of the colon or

XX

PT prostate, comprises 3351 human polynucleotide sequences.

XX

PS Claim 9; Page 562; 1046pp; English.

XX

CC the polynucleotide and for detection of transcription levels. Ribozymes  
 CC or antisense oligonucleotides can be generated. The polynucleotides and  
 CC their gene products are used as genetic or biochemical markers (e.g. in  
 CC blood or tissues) that will detect the earliest changes along the  
 CC carcinogenesis pathway and/or monitor the efficacy of therapies and  
 CC preventive interventions. The polynucleotides, polypeptides and  
 CC antibodies against them can be used in pharmaceutical compositions to  
 CC treat the cancers and proliferative disorders such as neoplasia,  
 CC dysplasia and hyperplasia  
 XX

SQ Sequence 400 BP; 123 A; 64 C; 119 G; 94 T; 0 U; 0 Other;

Alignment Scores:  
 Pred. No.: 17.2 Length: 400  
 Score: 51.00 Matches: 11  
 Percent Similarity: 70.00% Conservative: 3  
 Best Local Similarity: 55.00% Mismatches: 4  
 Query Match: 51.52% Indels: 2  
 DB: 5 Gaps: 1

US-10-799-747-116 (1-20) x AAF64369 (1-400)

QY 2 AlaAlaHisSerValLeuSer-----PheLeuLeuThrProTyrAlaLeuLysSer 19

Db 198 GCTCCCACTCTGTGCAAGTGATCATTCTTCATTTGACGCCCATTCGACTCTCTGCA 139

RESULT 11

AAH03292/c

ID AAH03292 standard; cDNA; 505 BP.

XX

AC AAH03292;

XX

DT 26-JUN-2001 (first entry)

XX

DE Human cDNA clone (5'-primer) SEQ ID NO:127.

XX

KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.

XX

OS Homo sapiens.

XX

PN EP1074617-A2.

XX

PD 07-FEB-2001.

XX

PF 28-JUL-2000; 2000EP-00116126.

XX

PR 29-JUL-1999; 99JP-00248036.

XX

PR 27-AUG-1999; 99JP-00300253.

XX

PR 11-JAN-2000; 2000JP-00118776.

XX

PR 02-MAY-2000; 2000JP-00183767.

XX

PR 09-JUN-2000; 2000JP-00241899.

XX

PA (HELI-) HELIX RES INST.

XX

PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;

XX

PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;

XX

DR WPI; 2001-318749/34.

XX

PT Primer sets for synthesizing polynucleotides, particularly the 5602 full-

XX

PT length cDNAs defined in the specification, and for the detection and/or

XX

PT diagnosis of the abnormality of the proteins encoded by the full-length

XX

XX cDNAs.

PS Claim 1; SEQ ID NO 127; 2537pp + Sequence Listing; English.

XX

CC The present invention describes primer sets for synthesizing 5602 full-

XX

CC length cDNAs defined in the specification. Where a primer set comprises:

XX

CC (a) an oligo-dr primer and an oligonucleotide complementary to the

XX

CC complementary strand of a polynucleotide which comprises one of the 5602

XX

CC nucleotide sequences defined in the specification, where the

XX

CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination

XX

CC of an oligonucleotide comprising a sequence complementary to the





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Alignment Scores:
Pred. No.:          90.6           Length:      1544
Score:              51.00          Matches:     10
Percent Similarity: 80.00%         Conservative: 2
Best Local Similarity: 66.67%       Mismatches:   3
Query Match:        51.52%         Indels:       0
DB:                  2             Gaps:         0

US-10-799-747-116 (1-20) x AA#86839 (1-1544)

QY    3  AlaHisSerValLeuSerPheIeuLeuTThrProTyrAlaLeu 17
      ||| :||::|||
Db    759 GCATTAACGGCTCTATTCTTCCTCTCGACCCTCAATG 803

```

RESULT 14  
AAH15239/c  
ID AAH15239 standard; cDNA; 1568 BP.  
XX  
XX  
XX AAH15239;  
XX AC  
XX  
26-JUN-2001 (first entry)  
DT  
DE Human cDNA sequence SEQ ID NO:13356.  
DE

Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.  
XX  
OS Homo sapiens.

XX	PN	EP1074617-A2.
XX	XX	
XX	PD	07-FEB-2001.
XX	XX	
XX	PF	28-JUL-2000; 2000EP-00116136.
XX	XX	
XX	PR	29-JUL-1999; 99JP-00248036.
XX	PR	27-AUG-1999; 99JP-00300253.
XX	PR	11-JAN-2000; 2000JP-00118776.
XX	PR	02-MAY-2000; 2000JP-00183767.
XX	PR	09-JUN-2000; 2000JP-00241899.
XX	XX	
XX	PA	(HELI-) HELIX RES INST.

Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;  
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
XX  
DR WPI; 2001-318749/34

Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs.

PS  
v  
Claim 8; SEQ ID NO 13356; 2537pp + Sequence Listing; English.

The present invention describes primer sets for synthesising 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dT primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesising polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893

CC represent human amino acid sequences; and AAH13629 to AAH13632 represent  
CC oligonucleotides, all of which are used in the exemplification of the  
CC present invention  
XX  
SQ Sequence 1568 BP; 447 A; 306 C; 425 G; 390 T; 0 U; 0 Other;

Alignment Scores:			
Pred. No.:	92.4	Length:	1568
Score:	51.00	Matches:	118
Percent Similarity:	70.00%	Conservative:	3
Best Local Similarity:	55.00%	Mismatches:	4
Query Match:	51.52%	Indels:	2
DB:	4	Gaps:	1

US-10-799-747-116 (1-20) x AAH15239 (1-1568)

Qy 2 AlaAlaHisSerValLeuSer-----PheLeuLeuTrpThrProTyrAlaLeuLysSer 19  
 |||:::||||| ||| |||||:::||||| |||||  
 Db 132 GCCTCCCACTCTTGTCGAAGTGATCACTTCCTCATTTGGACGCCCATTCACCTCTCTGCA 73

RESULT 15

AAH77480/C

ID AAH77480 standard; DNA; 2500 bp.

AA  
AC  
AAH77480:

DT 20-NOV-2001 (first entry)

XX

DE Mandrill immunodeficiency virus SIM27 related coding sequence #4.  
yy

Simian immunodeficiency virus 077

antibody detection: ds

XX  
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33

OS Simian immunod

XX  
PN  
NQ200003889-X

XX  
XX  
-E99C00007017

PD 05-FEB-2001.

XXV

28-JUL-2000; 2000NO-00003889.  
PF  
YY

03-AUG-1999. 99DE-01026003

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XXXX-0000000000

PA (DADE-) DADE BEHRING MARBURG GMBH.

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PI  
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Guertler LG, Kaptue

WPI; 2001-201061/20.

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PT  
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PT

PS Claim 1; Page 11-12; 47pp; Norwegian.

The present invention relates to a new mandrill immunodeficiency virus SIM27, which can be used to detect antibodies directed against the virus. The present sequence is a coding sequence described in the exemplification of the invention

Sequence 2500 BP; 872 A; 498 C; 637 G; 483 T; 0 U; 10 Other:

**Alignment Scores:**

Pred. No.:	164	Length:	2500
Score:	51.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	3
Local Similarity:	72.73%	Mismatches:	0
Query Match:	51.52%	Indels:	0
DB:	4	Gaps:	0

US-10-799-747-116 (1-20) x AAH77480 (1-2500)

7 LeuSerPheLeuLeuTrpThrProTyrAlaLeu 17

us-10-799-747-116.rng

Wed Jul 21 09:11:30 2004

Db 1453 TTGAGCTTTATACCTTTGGTCCCTACTCCTTG 1421

Search completed: July 21, 2004, 01:43:18  
Job time : 356 secs

GenCore version 5.1.6  
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 01:24:25 ; Search time 72 Seconds

(without alignments)

154.153 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 99

Sequence: 1 MAHSVLSFLLTTPYALKSX 20

Scoring table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5

Ygapop 10.0 , Ygapext 0.5

Fgapop 6.0 , Fgapext 7.0

Delop 6.0 , Delext 7.0

Searched: 682709 seqs, 277475446 residues

Total number of hits satisfying chosen parameters: 1365418

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters: -DEV=xlp

-O=/cgn2\_1/USPTO.spool.p/US10799747/runat.19072004.161418.21467/app.query.fasta\_1.199

-DB=Issued Patents NA -QFMT=fastap -SUFFIX=rni -MINMATCH=0.1 -LOOPCL=0

-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human0.cdi

-LIST=45 -DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15

-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000

-USER=US10799747 @CGN 1.1.105 @runat.19072004.161418.21467 -NCPU=6 -ICPU=3

-NO WMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPLOCK=100 -LONGLOG

-DRV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6

-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : Issued Patents NA:\*

1: /cgn2\_6/prodata/2/ina/5A.COMB.seq:\*

2: /cgn2\_6/prodata/2/ina/5B.COMB.seq:\*

3: /cgn2\_6/prodata/2/ina/6A.COMB.seq:\*

4: /cgn2\_6/prodata/2/ina/6B.COMB.seq:\*

5: /cgn2\_6/prodata/2/ina/PCTUS.COMB.seq:\*

6: /cgn2\_6/prodata/2/ina/backfiles1.seq:\*

pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	51	51.5	2500	4	US-09-625-972-20
C 2	51	51.5	9641	4	US-09-625-972-22
C 3	49	49.5	1368	4	US-09-134-000C-1177
C 4	48	48.5	207	4	US-09-313-294A-7069
C 5	48	48.5	1601	3	US-08-492-459-17
C 6	48	48.5	1601	3	US-08-423-752-17
C 7	48	48.5	1601	3	US-08-423-752-17
C 8	48	48.5	1601	3	US-08-716-873-31
C 9	48	48.5	1601	3	US-08-716-873-31
C 10	48	48.5	1601	3	US-09-368-431-31
C 11	48	48.5	1601	3	US-09-368-431-31
C 12	48	48.5	1601	3	US-09-368-431-32

C 13	48	48.5	1601	4	US-09-414-006-17	Sequence 17, Appl
C 14	48	48.5	1601	4	US-09-414-006-18	Sequence 18, Appl
C 15	48	48.5	1601	4	US-09-447-223-17	Sequence 17, Appl
C 16	48	48.5	1601	4	US-09-447-223-18	Sequence 18, Appl
17	48	48.5	2385	3	US-08-492-459-1	Sequence 1, Appl
18	48	48.5	2385	3	US-08-492-459-3	Sequence 3, Appl
19	48	48.5	2385	3	US-08-423-752-1	Sequence 1, Appl
20	48	48.5	2385	3	US-08-423-752-3	Sequence 3, Appl
21	48	48.5	2385	3	US-08-716-873-15	Sequence 15, Appl
22	48	48.5	2385	3	US-08-716-873-17	Sequence 17, Appl
23	48	48.5	2385	3	US-09-368-431-15	Sequence 15, Appl
24	48	48.5	2385	3	US-09-368-431-17	Sequence 17, Appl
25	48	48.5	2385	4	US-09-414-006-1	Sequence 1, Appl
26	48	48.5	2385	4	US-09-414-006-3	Sequence 3, Appl
27	48	48.5	2385	4	US-09-447-223-1	Sequence 1, Appl
28	48	48.5	2385	4	US-09-447-223-3	Sequence 3, Appl
C 29	48	48.5	17000	4	US-09-679-299A-18	Sequence 18, Appl
30	47	47.5	2210	4	US-09-016-434-1177	Sequence 1177, Ap
31	46	46.5	292	4	US-09-313-294A-4459	Sequence 4459, Ap
32	46	46.5	1098	4	US-09-718-692-3	Sequence 3, Appl
33	46	46.5	1098	4	US-09-718-852-3	Sequence 3, Appl
34	46	46.5	1098	4	US-09-718-815-3	Sequence 3, Appl
C 35	46	46.5	1296	4	US-09-328-352-467	Sequence 467, App
36	46	46.5	2595	4	US-09-016-434-1178	Sequence 1178, Ap
37	46	46.5	4911	4	US-09-718-692-1	Sequence 1, Appl
38	46	46.5	4911	4	US-09-718-852-1	Sequence 1, Appl
39	46	46.5	4911	4	US-09-718-815-1	Sequence 1, Appl
C 40	45.5	46.0	1230025	4	US-09-198-452A-1	Sequence 1, Appl
41	45	45.5	1495	4	US-09-016-434-1190	Sequence 1190, Ap
42	45	45.5	1495	4	US-09-023-655-1021	Sequence 1021, Ap
C 43	45	45.5	1896	4	US-09-134-000C-616	Sequence 616, App
44	45	45.5	2156	1	US-08-012-988A-1	Sequence 1, Appl
45	45	45.5	2156	4	US-09-023-655-1247	Sequence 1247, Ap

#### ALIGNMENTS

#### RESULT 1

US-09-625-972-20/c

; Sequence 20, Application US/09625972

; Patent No. 6566513

; GENERAL INFORMATION:

; APPLICANT: GUERTLER, Lutz Gerhard

; APPLICANT: HAUSER, Hans Peter

; APPLICANT: DONGMO DELOKO, Yvette Beatrice

; APPLICANT: ZEKENG, Leopold

; APPLICANT: KAPTUE, Lazare

; TITLE OF INVENTION: LENTIVIRUS FROM THE GROUP OF IMMUNODEFICIENCY VIRUSES OF DRILL M

; TITLE OF INVENTION: (MANDRILLUS LEUCOPHAUS) AND THEIR USE

; FILE REFERENCE: 067595/0106

; CURRENT APPLICATION NUMBER: US/09/625,972

; CURRENT FILING DATE: 2000-07-29

; PRIOR APPLICATION NUMBER: DE 199 36 003.0

; PRIOR FILING DATE: 1999-08-03

; NUMBER OF SEQ ID NOS: 57

; SOFTWARE: PatentIn version 3.0

; SEQ ID NO 20

; LENGTH: 2500

; TYPE: DNA

; ORGANISM: SIV - viral

US-09-625-972-20

Alignment Scores:	36.3	Length:	2500
Pred. No.:	51.00	Matches:	8
Score:	100.00%	Conservative:	3
Percent Similarity:	72.73%	Mismatches:	0
Best Local Similarity:	72.73%	Indels:	0
Query Match:	51.52%	Gaps:	0

US-10-799-747-116 (1-20) x US-09-625-972-20 (1-2500)

QY 7 LeuSerPheLeuLeuTrpThrProTyrAlaLeu 17



```
; TELEPHONE: 202-371-8850
; TELEFAX:
; TELEX:
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1601
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; ANTI-SENSE: Yes
US-08-492-459-17

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-492-459-17 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17
|||||:|||||
Db 843 CATAGCATCCTTGATATTITGGCCTGGTTCATATGGAGTC 802

RESULT 6
US-08-492-459-18/c
; Sequence 18, Application US/08492459
; Patent No. 6015689
; GENERAL INFORMATION:
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: REGULATION OF AUROBASIDIN SENSITIVITY IN FUNGUS
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wenderoth, Lind & Ponack
; STREET: 805 Fifteenth Street, N.W., #700
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.4 mb
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: Wordperfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/492,459
; FILING DATE: June 20, 1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/243,403
; FILING DATE: May 16, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Warren M. Cheek, Jr.
; REGISTRATION NUMBER: 33,367
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-8850
; TELEFAX:
; TELEX:
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1601
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: mRNA
; ANTI-SENSE: Yes
US-08-492-459-18

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-492-459-17 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17
|||||:|||||
Db 843 CATAGCATCCTTGATATTITGGCCTGGTTCATATGGAGTC 802

RESULT 7
US-08-423-752-17/c
; Sequence 17, Application US/08423752
; Patent No. 6022949
; GENERAL INFORMATION:
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: A GENE CODING FOR A PROTEIN REGULATING
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wenderoth, Lind & Ponack
; STREET: 805 Fifteenth Street, N.W., #700
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: Wordperfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/423,752
; FILING DATE: April 18, 1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/243,403
; FILING DATE: May 16, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Warren M. Cheek, Jr.
; REGISTRATION NUMBER: 33,367
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-371-8850
; TELEFAX:
; TELEX:
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1601
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; ANTI-SENSE: Yes
US-08-423-752-17

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-423-752-17 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17
|||||:|||||
Db 843 CATAGCATCCTTGATATTITGGCCTGGTTCATATGGAGTC 802
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RESULT 8  
US-08-423-752-18/c  
; Sequence 18, Application US/08423752  
; Patent No. 6022349  
; GENERAL INFORMATION:  
; APPLICANT: Takashi OKADO et al.  
; TITLE OF INVENTION: A GENE CODING FOR A PROTEIN REGULATING  
; TITLE OF INVENTION: AUROBASIDIN SENSITIVITY  
; NUMBER OF SEQUENCES: 22  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Wenderoth, Lind & Ponack  
; STREET: 805 Fifteenth Street, N.W., #700  
; CITY: Washington  
; STATE: D.C.  
; COUNTRY: U.S.A.  
; ZIP: 20005  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 5.25 inch, 500 kb  
; COMPUTER: IBM Compatible  
; OPERATING SYSTEM: MS-DOS  
; SOFTWARE: Wordperfect 5.1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/423,752  
; FILING DATE: April 18, 1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/243,403  
; FILING DATE: May 16, 1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Warren M. Cheek, Jr.  
; REGISTRATION NUMBER: 33,367  
; REFERENCE/DOCKET NUMBER:  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 202-371-8850  
; TELEFAX:  
; TELEX:  
; INFORMATION FOR SEQ ID NO: 18:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 1601  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: mRNA  
; ANTI-SENSE: Yes  
; US-08-423-752-18  
Alignment Scores:  
Pred. No.: 68.2 Length: 1601  
Score: 48.00 Matches: 7  
Percent Similarity: 64.29% Conservative: 2  
Best Local Similarity: 50.00% Mismatches: 5  
Query Match: 48.48% Indels: 0  
DB: 3 Gaps: 0  
US-10-799-747-116 (1-20) x US-08-423-752-18 (1-1601)  
Qy 4 HisSerValLeuSerPheLeuThrProTyrAlaLeu 17  
Db 843 CATAGCATCCTTGATATTGGCTGGTCCATATGGAGTC 802  
RESULT 9  
US-08-716-873-31/c  
; Sequence 31, Application US/08716873  
; Patent No. 6194166  
; GENERAL INFORMATION:  
; APPLICANT: Takashi OKADO et al.  
; TITLE OF INVENTION: GENE REGULATING AUROBASIDIN SENSITIVITY  
; NUMBER OF SEQUENCES: 50  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Wenderoth, Lind & Ponack  
; STREET: 805 Fifteenth Street, N.W., #700  
; CITY: Washington  
; STATE: D.C.  
; COUNTRY: U.S.A.  
; ZIP: 20005  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 5.25 inch, 500 kb  
; COMPUTER: IBM Compatible  
; OPERATING SYSTEM: MS-DOS  
; SOFTWARE: Wordperfect 5.1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/423,752  
; FILING DATE: April 18, 1995  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: 08/243,403  
; FILING DATE: May 16, 1994  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Warren M. Cheek, Jr.  
; REGISTRATION NUMBER: 33,367  
; REFERENCE/DOCKET NUMBER:  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 202-371-8850  
; TELEFAX:  
; TELEX:  
; INFORMATION FOR SEQ ID NO: 18:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 1601  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: mRNA  
; ANTI-SENSE: Yes  
; US-08-423-752-18  
Alignment Scores:  
Pred. No.: 68.2 Length: 1601  
Score: 48.00 Matches: 7  
Percent Similarity: 64.29% Conservative: 2  
Best Local Similarity: 50.00% Mismatches: 5  
Query Match: 48.48% Indels: 0  
DB: 3 Gaps: 0  
US-10-799-747-116 (1-20) x US-08-423-752-18 (1-1601)  
Qy 4 HisSerValLeuSerPheLeuThrProTyrAlaLeu 17  
Db 843 CATAGCATCCTTGATATTGGCTGGTCCATATGGAGTC 802

; COUNTRY: U.S.A.  
; ZIP: 20005  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 mb  
; COMPUTER: IBM Compatible  
; OPERATING SYSTEM: MS-DOS  
; SOFTWARE: Wordperfect 5.1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/716,873  
; FILING DATE: September 20, 1996  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER:  
; FILING DATE:  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Warren M. Cheek, Jr.  
; REGISTRATION NUMBER: 33,367  
; REFERENCE/DOCKET NUMBER:  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: 202-371-8850  
; TELEFAX:  
; TELEX:  
; INFORMATION FOR SEQ ID NO: 31:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 1601 bases  
; TYPE: nucleic acid  
; STRANDEDNESS: double  
; TOPOLOGY: linear  
; MOLECULE TYPE: genomic DNA  
; ANTI-SENSE: Yes  
; US-08-716-873-31  
Alignment Scores:  
Pred. No.: 68.2 Length: 1601  
Score: 48.00 Matches: 7  
Percent Similarity: 64.29% Conservative: 2  
Best Local Similarity: 50.00% Mismatches: 5  
Query Match: 48.48% Indels: 0  
DB: 3 Gaps: 0  
US-10-799-747-116 (1-20) x US-08-716-873-31 (1-1601)  
Qy 4 HisSerValLeuSerPheLeuThrProTyrAlaLeu 17  
Db 843 CATAGCATCCTTGATATTGGCTGGTCCATATGGAGTC 802  
RESULT 10  
US-08-716-873-32/c  
; Sequence 32, Application US/08716873  
; Patent No. 6194166  
; GENERAL INFORMATION:  
; APPLICANT: Takashi OKADO et al.  
; TITLE OF INVENTION: GENE REGULATING AUROBASIDIN SENSITIVITY  
; NUMBER OF SEQUENCES: 50  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Wenderoth, Lind & Ponack  
; STREET: 805 Fifteenth Street, N.W., #700  
; CITY: Washington  
; STATE: D.C.  
; COUNTRY: U.S.A.  
; ZIP: 20005  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Diskette, 3.5 inch, 1.44 mb  
; COMPUTER: IBM Compatible  
; OPERATING SYSTEM: MS-DOS  
; SOFTWARE: Wordperfect 5.1  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/716,873  
; FILING DATE: September 20, 1996  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER:  
; FILING DATE:

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/ ATTORNEY/AGENT INFORMATION:
/ NAME: Warren M. Cheek, Jr.
/ REGISTRATION NUMBER: 33,367
/ REFERENCE/DOCKET NUMBER:
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 202-371-8850
/ TELEFAX:
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 32:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 1601 bases
/ TYPE: nucleic acid
/ STRANDEDNESS: double
/ TOPOLOGY: linear
/ MOLECULE TYPE: mRNA
/ ANTI-SENSE: Yes
/ US-08-716-873-32

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-716-873-32 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17
|||||:::|||||
Db 843 CATAGCATCTTGATATTGGCTGGTTCATATGGAGTC 802

RESULT 11
US-09-368-431-31/c
/ Sequence 31, Application US/09368431
/ Patent No. 6294651
/ GENERAL INFORMATION:
/ APPLICANT: Takashi OKADO et al.
/ TITLE OF INVENTION: GENE REGULATING AUREOBASIDIN SENSITIVITY
/ TITLE OF INVENTION: (AS AMENDED)
/ NUMBER OF SEQUENCES: 50
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: WENDEROTH, LIND & PONACK, L.L.P.
/ STREET: 2033 K Street, N.W., #800
/ CITY: Washington
/ STATE: D.C.
/ COUNTRY: U.S.A.
/ ZIP: 20006
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Diskette, 3.5 inch, 1.44 mb
/ COMPUTER: IBM Compatible
/ OPERATING SYSTEM: MS-DOS
/ SOFTWARE: Wordperfect 5.1
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/368,431
/ FILING DATE: August 5, 1999
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 08/716,873
/ FILING DATE: September 20, 1996
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Warren M. Cheek, Jr.
/ REGISTRATION NUMBER: 33,367
/ REFERENCE/DOCKET NUMBER:
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 202-721-8200
/ TELEFAX: 202-721-8250
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 31:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 1601 bases
/ TYPE: nucleic acid
/ STRANDEDNESS: double
```

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/ TOPOLOGY: linear
/ MOLECULE TYPE: genomic DNA
/ ANTI-SENSE: Yes
/ US-09-368-431-31

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-09-368-431-31 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTTPThrProTyrAlaLeu 17
|||||:::|||||
Db 843 CATAGCATCTTGATATTGGCTGGTTCATATGGAGTC 802

RESULT 12
US-09-368-431-32/c
/ Sequence 32, Application US/09368431
/ Patent No. 6294651
/ GENERAL INFORMATION:
/ APPLICANT: Takashi OKADO et al.
/ TITLE OF INVENTION: GENE REGULATING AUREOBASIDIN SENSITIVITY
/ TITLE OF INVENTION: (AS AMENDED)
/ NUMBER OF SEQUENCES: 50
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: WENDEROTH, LIND & PONACK, L.L.P.
/ STREET: 2033 K Street, N.W., #800
/ CITY: Washington
/ STATE: D.C.
/ COUNTRY: U.S.A.
/ ZIP: 20006
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Diskette, 3.5 inch, 1.44 mb
/ COMPUTER: IBM Compatible
/ OPERATING SYSTEM: MS-DOS
/ SOFTWARE: Wordperfect 5.1
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/368,431
/ FILING DATE: August 5, 1999
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 08/716,873
/ FILING DATE: September 20, 1996
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Warren M. Cheek, Jr.
/ REGISTRATION NUMBER: 33,367
/ REFERENCE/DOCKET NUMBER:
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 202-721-8200
/ TELEFAX: 202-721-8250
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 32:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 1601 bases
/ TYPE: nucleic acid
/ STRANDEDNESS: double
/ TOPOLOGY: linear
/ MOLECULE TYPE: mRNA
/ ANTI-SENSE: Yes
/ US-09-368-431-32

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 3 Gaps: 0
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US-10-799-747-116 (1-20) x US-09-368-431-32 (1-1601)
;
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: REGULATION OF AUREOBASIDIN SENSITIVITY (AS
; AMENDED)
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Wenderoth, Lind & Ponack, L.L.P.
; STREET: 2033 K Street, N.W., #800
; CITY: Washington
; STATE: D.C.
; COUNTRY: U.S.A.
; ZIP: 20006
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.5 inch, 1.4 mb
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: MS-DOS
; SOFTWARE: Wordperfect 5.1
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/414,006
; FILING DATE: October 7, 1999
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/492,459
; FILING DATE: June 20, 1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/243,403
; FILING DATE: May 16, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Warren M. Cheek, Jr.
; REGISTRATION NUMBER: 33,367
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-721-8200
; TELEFAX: 202-721-8250
; TELEX:
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1601
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: genomic DNA
; ANTI-SENSE: Yes
; US-09-414-006-17
;
; Alignment Scores:
; Pred. No.: 68.2 Length: 1601
; Score: 48.00 Matches: 7
; Percent Similarity: 64.29% Conservative: 2
; Best Local Similarity: 50.00% Mismatches: 5
; Query Match: 48.48% Indels: 0
; DB: 4 Gaps: 0
;
; US-10-799-747-116 (1-20) x US-09-414-006-18 (1-1601)
;
; Qy 4 HisSerValLeuSerPheLeuLeuTrpThrProTyrAlaLeu 17
; Db 843 CATAGCATCCTTGATATTTCGCTGGGTTCCATATGGAGTC 802
;
; RESULT 15
; US-09-447-223-17/c
; Sequence 17, Application US/09447223
; Patent No. 6432664
; GENERAL INFORMATION:
; APPLICANT: Takashi OKADO et al.
; TITLE OF INVENTION: A GENE CODING FOR A PROTEIN REGULATING
; CORRESPONDENCE ADDRESS:
; NUMBER OF SEQUENCES: 22
; ADDRESSEE: Wenderoth, Lind & Ponack
; STREET: 805 Fifteenth Street, N.W., #700
; CITY: Washington
; STATE: D.C.
;
; US-10-799-747-116 (1-20) x US-09-414-006-17 (1-1601)
;
; Qy 4 HisSerValLeuSerPheLeuLeuTrpThrProTyrAlaLeu 17
; Db 843 CATAGCATCCTTGATATTTCGCTGGGTTCCATATGGAGTC 802
;
; RESULT 14
; US-09-414-006-18/c
; Sequence 18, Application US/09414006
; Patent No. 6348577
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: COUNTRY: U.S.A.
: ZIP: 20005
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette, 5.25 inch, 500 kb
: COMPUTER: IBM Compatible
: OPERATING SYSTEM: MS-DOS
: SOFTWARE: Wordperfect 5.1
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: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/09/447,223
: FILING DATE: 23-No. 6432664-1999
: CLASSIFICATION: <Unknown>
:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: 08/243,403
: FILING DATE: May 16, 1994
:
: ATTORNEY/AGENT INFORMATION:
: NAME: Warren M. Cheek, Jr.
: REGISTRATION NUMBER: 33,367
: REFERENCE/DOCKET NUMBER: <Unknown>
:
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 202-371-8850
: TELEFAX: <Unknown>
: TELEX: <Unknown>
:
: INFORMATION FOR SEQ ID NO: 17:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1601
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: genomic DNA
: ANTI-SENSE: Yes
: SEQUENCE DESCRIPTION: SEQ ID NO: 17:
US-09-447-223-17

Alignment Scores:
Pred. No.: 68.2 Length: 1601
Score: 48.00 Matches: 7
Percent Similarity: 64.29% Conservative: 2
Best Local Similarity: 50.00% Mismatches: 5
Query Match: 48.48% Indels: 0
DB: 4 Gaps: 0

US-10-799-747-116 (1-20) X US-09-447-223-17 (1-1601)

QY 4 HisSerValLeuSerPheLeuLeuTrpThrProTyrAlaLeu 17
Db 843 CATAGCATCCTGTGATATTGGCTGGTTCATATGGAGTC 802

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Search completed: July 21, 2004, 02:54:35  
Job time : 75 secs

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QM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 02:14:01 ; Search time 355 Seconds  
(without alignments)  
274.852 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 99

Sequence: 1 MAHSLVSLFWLTPYALKSX 20

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Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 3191023 seqs, 2439312756 residues

Total number of hits satisfying chosen parameters: 6382046

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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-FCAPOP=6 -FCAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

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16: /cgn2\_6/ptodata/2/pubpna/US10C\_PUBCOMB.seq:  
17: /cgn2\_6/ptodata/2/pubpna/US10\_NEW\_PUB.seq:  
18: /cgn2\_6/ptodata/2/pubpna/US60\_NEW\_PUB.seq:  
19: /cgn2\_6/ptodata/2/pubpna/US60\_PUBCOMB.seq:

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Match	Length	DB ID	Description
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1	98	99.0	1434	15	US-10-195-730-12	Sequence 12, Appl
2	57	57.6	87731	13	US-10-087-192-1342	Sequence 1342, Ap
3	56	56.6	8417	16	US-10-074-024-635	Sequence 635, App
4	53	53.5	5760	16	US-10-074-024-426	Sequence 426, App
5	53	53.5	5763	16	US-10-074-024-425	Sequence 425, App
6	51	51.5	2500	15	US-10-364-360-20	Sequence 20, Appl
7	51	51.5	9641	15	US-10-364-360-22	Sequence 22, Appl
8	51	51.5	56577	13	US-10-087-192-1396	Sequence 1396, Ap
9	51	51.5	2940917	13	US-10-027-632-174763	Sequence 174763, Ap
10	51	51.5	2940917	16	US-10-027-632-174763	Sequence 174763, Ap
11	50	50.5	620	12	US-10-152-319A-215	Sequence 215, App
12	50	50.5	740	13	US-10-027-632-11891	Sequence 11891, A
13	50	50.5	740	16	US-10-027-632-11891	Sequence 11891, A
14	50	50.5	1206	13	US-10-027-632-250731	Sequence 250731, A
15	50	50.5	1206	13	US-10-027-632-250731	Sequence 250731, A
16	50	50.5	1206	16	US-10-027-632-250731	Sequence 250732, A
17	50	50.5	1206	16	US-10-027-632-250731	Sequence 250732, A
18	50	50.5	3034	17	US-10-437-963-7778	Sequence 7778, Ap
19	49	49.5	519	13	US-10-027-632-284792	Sequence 284792, A
20	49	49.5	519	16	US-10-027-632-284792	Sequence 284792, A
21	49	49.5	793	13	US-10-027-632-144194	Sequence 144194, A
22	49	49.5	793	13	US-10-027-632-144194	Sequence 144194, A
23	49	49.5	793	16	US-10-027-632-144194	Sequence 144195, A
24	49	49.5	793	16	US-10-027-632-144194	Sequence 144195, A
25	49	49.5	808	13	US-10-027-632-155999	Sequence 155999, A
26	49	49.5	808	16	US-10-027-632-155999	Sequence 155999, A
27	49	49.5	828	13	US-10-027-632-158555	Sequence 158555, A
28	49	49.5	828	13	US-10-027-632-158555	Sequence 158555, A
29	49	49.5	828	16	US-10-027-632-158555	Sequence 158555, A
30	49	49.5	828	16	US-10-027-632-158555	Sequence 158555, A
31	49	49.5	1092	17	US-10-437-963-61207	Sequence 61207, A
32	49	49.5	2393	13	US-10-027-632-102504	Sequence 102504, A
33	49	49.5	2393	16	US-10-027-632-102504	Sequence 102504, A
34	49	49.5	32768	9	US-09-070-927A-128	Sequence 128, App
35	48.5	49.0	72705	13	US-10-087-192-1966	Sequence 1966, Ap
36	48	48.5	290	14	US-10-040-739-637	Sequence 637, App
37	48	48.5	375	10	US-09-803-719-2326	Sequence 2326, Ap
38	48	48.5	495	12	US-09-969-034-1082	Sequence 1082, Ap
39	48	48.5	587	13	US-10-424-599-90540	Sequence 90540, A
40	48	48.5	612	14	US-10-002-344A-83	Sequence 83, Appl
41	48	48.5	802	13	US-10-424-599-111198	Sequence 111198, A
42	48	48.5	940	15	US-10-017-161-1223	Sequence 1223, Ap
43	48	48.5	940	16	US-10-292-798-1027	Sequence 1027, Ap
44	48	48.5	960	9	US-09-886-055-148	Sequence 148, App
45	48	48.5	960	10	US-09-804-291-148	Sequence 148, App

#### ALIGNMENTS

RESULT 1  
US-10-195-730-12  
; Sequence 12, Application US/10195730  
; Publication No. US20030144492A1  
; GENERAL INFORMATION:  
; APPLICANT: Rosen et. al  
; TITLE OF INVENTION: 101 Human Secreted Proteins  
; FILE REFERENCE: P2017P1  
; CURRENT APPLICATION NUMBER: US/10/195,730  
; CURRENT FILING DATE: 2002-07-16  
; PRIOR APPLICATION NUMBER: US/09/281,976  
; PRIOR FILING DATE: 1999-03-31  
; PRIOR APPLICATION NUMBER: 60/060,837  
; PRIOR FILING DATE: 1997-10-02  
; PRIOR APPLICATION NUMBER: 60/060,862  
; PRIOR FILING DATE: 1997-10-02  
; NUMBER OF SEQ ID NOS: 390  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 12  
; LENGTH: 1434  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-195-730-12

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; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (2140)
; OTHER INFORMATION: n equals a,t,g, or c
US-10-074-024-635
Alignment Scores:
Pred. No.: 75.7 Length: 8417
Score: 56.00 Matches: 9
Percent Similarity: 78.57% Conservatives: 2
Best Local Similarity: 64.29% Mismatches: 3
Query Match: 56.57% Indels: 0
DB: 16 Gaps: 0

US-10-799-747-116 (1-20) x US-10-074-024-635 (1-8417)
Qy 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTyrProTyrAlaLeu 14
Db 934 ATGCCACACATTCGATTGACGTTTTTATTCGGACCCCA 975

RESULT 4
US-10-074-024-426
; Sequence 426, Application US/10074024
; Publication No. US20030232975A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC001C1
; CURRENT APPLICATION NUMBER: US/10/074,024
; CURRENT FILING DATE: 2002-02-14
; Prior Application removed - See file Wrapper or Palm
; NUMBER OF SEQ ID NOS: 879
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 426
; LENGTH: 5760
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-074-024-426
Alignment Scores:
Pred. No.: 159 Length: 5760
Score: 53.00 Matches: 10
Percent Similarity: 68.75% Conservatives: 1
Best Local Similarity: 62.50% Mismatches: 5
Query Match: 53.54% Indels: 0
DB: 16 Gaps: 0

US-10-799-747-116 (1-20) x US-10-074-024-426 (1-5760)
Qy 2 AlaAlaHisSerValLeuSerPheLeuLeuTyrProTyrAlaLeu 17
Db 567 GCCCGGCACTCACCCCTTGCTTTGGCTTGTATATTCCTT 614

RESULT 5
US-10-074-024-425
; Sequence 425, Application US/10074024
; Publication No. US20030232975A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC001C1
; CURRENT APPLICATION NUMBER: US/10/074,024
; CURRENT FILING DATE: 2002-02-14
; Prior Application removed - See file Wrapper or Palm
; NUMBER OF SEQ ID NOS: 879
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 425
; LENGTH: 5763
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1871)
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; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (2140)
; OTHER INFORMATION: n equals a,t,g, or c
US-10-074-024-635
Alignment Scores:
Pred. No.: 1434 Length: 1434
Score: 98.00 Matches: 19
Percent Similarity: 100.00% Conservatives: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 98.99% Indels: 0
DB: 15 Gaps: 0

US-10-799-747-116 (1-20) x US-10-195-730-12 (1-1434)
Qy 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTyrProTyrAlaLeuLysSer 19
Db 507 ATGCAGCCCATTCAGCTTCTTCTCTCGACCTTNGCTCGAATCA 563

RESULT 2
US-10-087-192-1342
; Sequence 1342, Application US/10087192
; Publication No. US20020182586A1
; GENERAL INFORMATION:
; APPLICANT: Morris, David W.
; APPLICANT: Engelhard, Eric K.
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR
; TITLE OF INVENTION: CANCER
; FILE REFERENCE: 529452000122
; CURRENT APPLICATION NUMBER: US/10/087,192
; CURRENT FILING DATE: 2002-03-01
; PRIOR APPLICATION NUMBER: US 09/747,377
; PRIOR FILING DATE: 2000-12-22
; PRIOR APPLICATION NUMBER: US 09/798,586
; PRIOR FILING DATE: 2001-03-02
; NUMBER OF SEQ ID NOS: 2059
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1342
; LENGTH: 87731
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-087-192-1342
Alignment Scores:
Pred. No.: 784 Length: 87731
Score: 57.00 Matches: 10
Percent Similarity: 81.25% Conservatives: 3
Best Local Similarity: 62.50% Mismatches: 3
Query Match: 57.58% Indels: 0
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-087-192-1342 (1-87731)
Qy 2 AlaAlaHisSerValLeuSerPheLeuLeuTyrProTyrAlaLeu 17
Db 41423 TCTAGTCATCCCATCTTGTCTTCTGAGCTGGACTCCTTACAGCTG 41470

RESULT 3
US-10-074-024-635
; Sequence 635, Application US/10074024
; Publication No. US20030232975A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC001C1
; CURRENT APPLICATION NUMBER: US/10/074,024
; CURRENT FILING DATE: 2002-02-14
; Prior Application removed - See file Wrapper or Palm
; NUMBER OF SEQ ID NOS: 879
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 635
; LENGTH: 8417
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (2138)
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; OTHER INFORMATION: n equals a,t,g, or c  
US-10-074-024-425

Alignment Scores: Length: 5763  
Pred. No.: 159 Matches: 10  
Score: 53.00  
Percent Similarity: 68.75% Conservative: 1  
Best Local Similarity: 62.50% Mismatches: 5  
Query Match: 53.54% Indels: 0  
DB: 16 Gaps: 0

US-10-799-747-116 (1-20) x US-10-074-024-425 (1-5763)

Qy 2 AlaAlaHisSerValLeuSerPheLeuLeuTrrProTyrAlaLeu 17  
Db 567 GCCGGCACTCAACCCCTTGCTTTGGCTTTGGTTGTATATCTCTT 614

## RESULT 6

US-10-364-360-20/c  
; Sequence 20, Application US/10364360  
; Publication No. US20030180324A1  
; GENERAL INFORMATION:  
; APPLICANT: GUERTLER, Lutz Gerhard  
; APPLICANT: HAUSER, Hans Peter  
; APPLICANT: DONGMO DELOKO, Yvette Beatrice  
; APPLICANT: ZEKENG, Leopold  
; APPLICANT: KAPTUE, Lazare  
; TITLE OF INVENTION: LENTIVIRUS FROM THE GROUP OF IMMUNODEFICIENCY VIRUSES OF DRILL MC  
; FILE OF INVENTION: (MANDRILLUS LEUCOPHAUS) AND THEIR USE  
; FILE REFERENCE: 067595/0106  
; CURRENT APPLICATION NUMBER: US/10/364,360  
; CURRENT FILING DATE: 2003-02-12  
; PRIOR APPLICATION NUMBER: US/09/625,972  
; PRIOR FILING DATE: 2000-07-29  
; PRIOR APPLICATION NUMBER: DE 199 36 003.0  
; PRIOR FILING DATE: 1999-08-03  
; NUMBER OF SEQ ID NOS: 57  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 20  
; LENGTH: 2500  
; TYPE: DNA  
; ORGANISM: SIV - viral  
US-10-364-360-20

Alignment Scores: Length: 2500  
Pred. No.: 133 Matches: 8  
Score: 51.00  
Percent Similarity: 100.00% Conservative: 3  
Best Local Similarity: 72.73% Mismatches: 0  
Query Match: 51.52% Indels: 0  
DB: 15 Gaps: 0

US-10-799-747-116 (1-20) x US-10-364-360-20 (1-2500)

Qy 7 LeuSerPheLeuLeuTrrProTyrAlaLeu 17  
Db 1453 TTGAGCTTTTACTTTGGTCCCTCTACTCCTTG 1421

## RESULT 7

US-10-364-360-22/c  
; Sequence 22, Application US/10364360  
; Publication No. US20030180324A1  
; GENERAL INFORMATION:  
; APPLICANT: GUERTLER, Lutz Gerhard  
; APPLICANT: HAUSER, Hans Peter  
; APPLICANT: DONGMO DELOKO, Yvette Beatrice  
; APPLICANT: ZEKENG, Leopold  
; APPLICANT: KAPTUE, Lazare  
; TITLE OF INVENTION: LENTIVIRUS FROM THE GROUP OF IMMUNODEFICIENCY VIRUSES OF DRILL MC  
; FILE OF INVENTION: (MANDRILLUS LEUCOPHAUS) AND THEIR USE  
; FILE REFERENCE: 067595/0106  
; CURRENT APPLICATION NUMBER: US/10/364,360  
; CURRENT FILING DATE: 2003-02-12

; PRIOR APPLICATION NUMBER: US/09/625,972  
; PRIOR FILING DATE: 2000-07-29  
; PRIOR APPLICATION NUMBER: DE 199 36 003.0  
; PRIOR FILING DATE: 1999-08-03  
; NUMBER OF SEQ ID NOS: 57  
; SOFTWARE: PatentIn version 3.0  
; SEQ ID NO 22  
; LENGTH: 9641  
; TYPE: DNA  
; ORGANISM: SIV - viral  
US-10-364-360-22

Alignment Scores: Length: 9641  
Pred. No.: 639 Matches: 8  
Score: 51.00  
Percent Similarity: 100.00% Conservative: 3  
Best Local Similarity: 72.73% Mismatches: 0  
Query Match: 51.52% Indels: 0  
DB: 15 Gaps: 0

US-10-799-747-116 (1-20) x US-10-364-360-22 (1-9641)

Qy 7 LeuSerPheLeuLeuTrrProTyrAlaLeu 17  
Db 1453 TTGAGCTTTTACTTTGGTCCCTCTACTCCTTG 1421

## RESULT 8

US-10-087-192-1396/c  
; Sequence 1396, Application US/10087192  
; Publication No. US20020182586A1  
; GENERAL INFORMATION:  
; APPLICANT: MORRIS, David W.  
; APPLICANT: ENGELHARD, Eric K.  
; TITLE OF INVENTION: NOVEL COMPOSITIONS AND METHODS FOR  
; TITLE OF INVENTION: CANCER  
; FILE REFERENCE: 529452000122  
; CURRENT APPLICATION NUMBER: US/10/087,192  
; CURRENT FILING DATE: 2002-03-01  
; PRIOR APPLICATION NUMBER: US 09/747,377  
; PRIOR FILING DATE: 2000-12-22  
; PRIOR APPLICATION NUMBER: US 09/798,586  
; PRIOR FILING DATE: 2001-03-02  
; NUMBER OF SEQ ID NOS: 2059  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 1396  
; LENGTH: 56577  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: (1)...(56577)  
; OTHER INFORMATION: n = A,T,C or G  
US-10-087-192-1396

Alignment Scores: Length: 56577  
Pred. No.: 5,03e+03 Matches: 10  
Score: 51.00  
Percent Similarity: 61.11% Conservative: 1  
Best Local Similarity: 55.56% Mismatches: 7  
Query Match: 51.52% Indels: 0  
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-087-192-1396 (1-56577)

Qy 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTrrProTyrAlaLeu 18  
Db 29892 ATGATAGCTCACTGCAGCTTGAACCTCTTGACTCCCAAGCTGCTGGAA 29839

## RESULT 9

US-10-027-632-174763/c  
; Sequence 174763, Application US/10027632  
; Publication No. US20020198371A1  
; GENERAL INFORMATION:

```
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 174763
; LENGTH: 2940917
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)..(2940917)
; OTHER INFORMATION: n = A,T,C or G
US-10-027-632-174763

Alignment Scores:
Pred. No.: 4.83e+05 Length: 2940917
Score: 51.00 Matches: 7
Percent Similarity: 84.62% Conservative: 4
Best Local Similarity: 53.85% Mismatches: 2
Query Match: 51.52% Indels: 0
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-174763 (1-2940917)

Qy 3 AlahisSerValLeuSerPheLeuLeuTrpThrProTyr 15
Db 189168 AGTCACCCATTTGTTTTGTTTGGACTCCATT 189130

RESULT 10
US-10-632-174763/c
; Sequence 174763, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; CURRENT FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0

US-10-027-632-174763

Alignment Scores:
Pred. No.: 4.83e+05 Length: 2940917
Score: 51.00 Matches: 7
Percent Similarity: 84.62% Conservative: 4
Best Local Similarity: 53.85% Mismatches: 2
Query Match: 51.52% Indels: 0
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-174763 (1-2940917)

Qy 3 AlahisSerValLeuSerPheLeuLeuTrpThrProTyr 15
Db 189168 AGTCACCCATTTGTTTTGTTTGGACTCCATT 189130

RESULT 11
US-10-152-319A-215/c
; Sequence 215, Application US/10152319A
; Publication No. US20040072160A1
; GENERAL INFORMATION:
; APPLICANT: Mendrick, Donna
; APPLICANT: Porter, Mark
; APPLICANT: Johnson, Kory
; APPLICANT: Higgs, Brandon
; APPLICANT: Castle, Arthur
; APPLICANT: Elashoff, Michael
; TITLE OF INVENTION: Molecular Toxicology Modeling
; FILE REFERENCE: 44921-5089-US
; CURRENT APPLICATION NUMBER: US/10/152,319A
; CURRENT FILING DATE: 2002-05-22
; PRIOR APPLICATION NUMBER: US 60/292,335
; PRIOR FILING DATE: 2001-05-22
; PRIOR APPLICATION NUMBER: US 60/297,523
; PRIOR FILING DATE: 2001-06-13
; PRIOR APPLICATION NUMBER: US 60/298,925
; PRIOR FILING DATE: 2001-06-19
; PRIOR APPLICATION NUMBER: US 60/303,810
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/303,807
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/303,808
; PRIOR FILING DATE: 2001-07-10
; PRIOR APPLICATION NUMBER: US 60/315,047
; PRIOR FILING DATE: 2001-08-28
; PRIOR APPLICATION NUMBER: US 60/324,928
; PRIOR FILING DATE: 2001-09-27
; PRIOR APPLICATION NUMBER: US 60/330,867
; PRIOR FILING DATE: 2001-11-01
; PRIOR APPLICATION NUMBER: US 60/330,462
; PRIOR FILING DATE: 2001-10-22
; Remaining Prior Application data removed - See File Wrapper or PALM.
; NUMBER OF SEQ ID NOS: 2221
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 215
; LENGTH: 620
; TYPE: DNA
; ORGANISM: Rattus norvegicus
; FEATURE:
; OTHER INFORMATION: Genbank Accession No. AA891812
; NAME/KEY: misc feature
; LOCATION: (1)..(620)
; OTHER INFORMATION: n = a or c or g or t
US-10-152-319A-215
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Best Local Similarity: 57.14% Mismatches: 4  
Query Match: 50.51% Indels: 0  
DB: 13 Gaps: 0  
US-10-799-747-116 (1-20) x US-10-027-632-250731 (1-1206)  
Qy 6 ValLeuSerPheLeuThrProTyrAlaLeuLysSer 19  
Db 704 ATTGTGACCTTCCTATTCTGGCTACCATACCCCATCTCCAGC 745

RESULT 15  
US-10-027-632-250732  
; Sequence 250732, Application US/10027632  
; Publication No. US20020198371A1  
; GENERAL INFORMATION:  
; APPLICANT: Wang, David G.  
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
; POLYMORPHISMS IN THE HUMAN GENOME  
; FILE REFERENCE: 108827.129  
; CURRENT APPLICATION NUMBER: US/10/027,632  
; CURRENT FILING DATE: 2002-04-30  
; PRIOR APPLICATION NUMBER: US 60/218,006  
; PRIOR FILING DATE: 2000-07-12  
; PRIOR APPLICATION NUMBER: US 60/198,676  
; PRIOR FILING DATE: 2000-04-20  
; PRIOR APPLICATION NUMBER: US 60/193,483  
; PRIOR FILING DATE: 2000-03-29  
; PRIOR APPLICATION NUMBER: US 60/185,218  
; PRIOR FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: US 60/167,363  
; PRIOR FILING DATE: 1999-11-23  
; PRIOR APPLICATION NUMBER: US 60/156,358  
; PRIOR FILING DATE: 1999-09-28  
; PRIOR APPLICATION NUMBER: US 60/146,002  
; PRIOR FILING DATE: 1999-08-09  
; NUMBER OF SEQ ID NOS: 325720  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 250732  
; LENGTH: 1206  
; TYPE: DNA  
; ORGANISM: Human  
US-10-027-632-250732

Alignment Scores:  
Pred. No.: 84.1 Length: 1206  
Score: 50.00 Matches: 8  
Percent Similarity: 71.43% Conservative: 2  
Best Local Similarity: 57.14% Mismatches: 4  
Query Match: 50.51% Indels: 0  
DB: 13 Gaps: 0  
US-10-799-747-116 (1-20) x US-10-027-632-250732 (1-1206)  
Qy 6 ValLeuSerPheLeuThrProTyrAlaLeuLysSer 19  
Db 704 ATTGTGACCTTCCTATTCTGGCTACCATACCCCATCTCCAGC 745

Search completed: July 21, 2004, 04:13:52  
Job time : 1132 secs



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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 01:17:25 ; Search time 2469 Seconds  
(without alignments)  
241.897 Million cell updates/sec

Title: US-10-799-747-116  
Perfect score: 99

Sequence: 1 MAHSVLSFLWTPYALKSX 20

Scoring table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5  
Ygapop 10.0 , Ygapext 0.5  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 27513289 seqs, 14931090276 residues

Total number of hits satisfying chosen parameters: 55026578

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlp  
-Q=/cgn2\_1/USPRO.spool.p/US10799747/runat 19072004 161416 21448/app\_query.fasta\_1.199  
-DB=EST -QPMT=fastcap -SUFFIX=rst -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0  
-UNITS=bits -START=1 -END=-1 -NATRIX=blosum62 -TRANS=human40.cdi -LIST=45  
-DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=200000000  
-USER=US10799747@cgn 1 15180 @runat 19072004 161416 21448 -NCPU=6 -ICPU=3  
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSBLOCK=100 -LONGLOG  
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6  
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

EST:  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_htc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_htc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rod:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrl:\*  
28: gb\_gss1:\*

29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	98	99.0	240	10	BF910533 CM4-UT004
C 2	98	99.0	575	12	BM722991 UI-E-R01-
C 3	98	99.0	765	12	BI914473 603182264
C 4	98	99.0	785	12	BG484396 602505037
C 5	98	99.0	1201	13	EX441923 BX441923
C 6	56	56.6	352	9	AA654972 nt61e09.s
C 7	55	55.6	642	29	CE789116 tigr-gss-
C 8	55	55.6	723	28	AZ831423 2M0111H12
C 9	54	54.5	550	29	TA226G11P
C 10	54	54.5	724	28	AQ645123 RPCI93-EC
C 11	54	54.5	816	14	CA967853 CCLX04a13
C 12	53	53.5	394	9	AA766276 nz38c04.s
C 13	53	53.5	429	9	AA804444
C 14	53	53.5	600	12	BI140807 IPI 39.B0
C 15	53	53.5	603	12	BM571235 fw75h06.x
C 16	53	53.5	635	13	BU262936 603818292
C 17	53	53.5	641	13	BU350638 603526071
C 18	53	53.5	641	13	BU448459 603211744
C 19	53	53.5	642	13	BU353991 603529626
C 20	53	53.5	643	13	BU394939 603511454
C 21	53	53.5	649	13	BU37269 603513967
C 22	53	53.5	657	13	BU39405 603512306
C 23	53	53.5	784	29	CG759218 ZMMBB031
C 24	53	53.5	804	28	AZ906788 RPCI-24-1
C 25	53	53.5	904	29	CG768437 TCB42.3 F
C 26	52.5	53.0	348	28	CC390782 PUH292TD
C 27	52	52.5	254	9	AA324608 EST27458
C 28	52	52.5	259	28	AQ104627 HS_3047_B
C 29	52	52.5	262	9	AI164098 A054p15u
C 30	52	52.5	288	14	F07932 HSC2LH031 n
C 31	52	52.5	293	10	BB564042 BB564042
C 32	52	52.5	372	29	CC717225 OGLCD72TV
C 33	52	52.5	374	9	AA411283 zv26f01.x
C 34	52	52.5	390	10	AW338878 ha58a11.x
C 35	52	52.5	423	28	CC370277 PUHGP23TD
C 36	52	52.5	584	9	AI754441 cr25d04.x
C 37	52	52.5	607	9	AI672090 ty63d04.x
C 38	52	52.5	653	28	CC004087 PUH064TD
C 39	52	52.5	669	28	AZ572182 302PVC03
C 40	52	52.5	691	13	BU871645 Q32H01 P
C 41	52	52.5	705	14	CD218841 pgrin.pk0
C 42	52	52.5	749	29	CG296747 OG3C11TV
C 43	52	52.5	759	12	BI837121 603090085
C 44	52	52.5	759	28	BZ985367 PUETC180TD
C 45	52	52.5	837	29	CC717215 OGLCD72TH

# ALIGNMENTS

RESULT 1  
BF910533/c  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS

BF910533  
CM4-UT0042-011100-395-f08 UT0042 Homo sapiens cDNA, mRNA sequence.  
BF910533  
BF910533.1 GI:12301991  
EST.  
Homo sapiens (human)  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R.,

240 bp mRNA linear EST 18-JAN-2001

**TITLE**  
Shotgun sequencing of the human transcriptome with ORF expressed  
sequence tags

**JOURNAL**  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)

**MEDLINE**  
20202663

**PUBMED**  
10737800

**COMMENT**  
Contact: Simpson A.J.G.  
Laboratory of Cancer Genetics  
Ludwig Institute for Cancer Research  
Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP,  
Brazil  
Tel: +55-11-2704922  
Fax: +55-11-2707001  
Email: asimpson@ludwig.org.br  
This sequence was derived from the FAPESP/LICR Human Cancer Genome  
Project. This entry can be seen in the following URL  
(http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM4&t2=CM4-UT0042-  
011100-395-f08&t3=2000-11-01&t4=1)  
Seq primer: puc 18 forward  
High quality sequence start: 81  
High quality sequence stop: 240.

**FEATURES**  
source  
1..240  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/dev\_stage="Adult"  
/clone\_lib="UT0042"  
/note="Organ: uterus\_tumor; Vector: puc18; Site 1: SmaI;  
Site 2: SmaI; A mini-library was made by cloning products  
derived from ORESTES PCR (U.S. Letters Patent application  
No. 196,716 - Ludwig Institute for Cancer Research)  
profiles into the pUC 18 vector. Reverse transcription of  
tissue mRNA and cDNA amplification were performed under  
low stringency conditions."

**ORIGIN**  
Alignment Scores:  
Pred. No.: 8 23e-06 Length: 240  
Score: 98.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 98.99% Indels: 0  
DB: 10 Gaps: 0

**ORIGIN**  
Alignment Scores:  
Pred. No.: 2 54e-05 Length: 575  
Score: 98.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 98.99% Indels: 0  
DB: 12 Gaps: 0

US-10-799-747-116 (1-20) x BF910533 (1-240)

QY 1 MetaAlaHisSerValLeuSerPheLeuThrProTyAlaLeuLysSer 19  
|||||  
DB 191 ATGGCAGCCCAATCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTTGAATCA 135

US-10-799-747-116 (1-20) x BM722991 (1-575)

QY 1 MetaAlaHisSerValLeuSerPheLeuThrProTyAlaLeuLysSer 19  
|||||  
DB 454 ATGGCAGCCCAATCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTTGAATCA 510

**RESULT 2**  
BM722991  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Homo sapiens (human)  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 575)  
Ronaldo, M.F., Lennon, G. and Soares, M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
Genome Res. 6 (9), 791-806 (1996)  
JOURNAL  
MEDLINE  
97044477

**RESULT 3**  
BI914473  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 765)  
NIH-MGC http://mgc.nci.nih.gov/.

**COMMENT**  
Contact: Soares, MB  
Coordinated Laboratory for Computational Genomics  
University of Iowa  
375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: bento-soares@uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa  
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa  
DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research  
Genetics (www.resgen.com).  
The following repetitive elements were found in this cDNA  
sequence: 148-169, >AT\_richLow\_complexity (matched complement)  
Seq primer: M13 Reverse.  
Location/Qualifiers  
1..575  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="UI-E-E01-aid-a-14-0-UI"  
/tissue\_type="fetal eye"  
/dev\_stage="fetal"  
/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"  
/clone\_lib="UI-E-E01"  
/note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a  
modified polylinker; Site 1: EcoR I; Site 2: Not I;  
UI-E-E01 is a normalized cDNA library containing the  
following tissue(s): fetal eye. The library was  
constructed according to Bonaldo, Lennon and Soares,  
Genome Research, 6:791-806, 1996. First strand cDNA  
synthesis was primed with an oligo-dT primer containing a  
Not I site. Double stranded cDNA was ligated to an EcoR I  
adaptor, digested with Not I, and cloned directionally  
into pT73-Pac vector. The oligonucleotide used to prime  
the synthesis of first-strand cDNA contains a library tag  
sequence that is located between the Not I site and the  
(dT)18 tail. The sequence tag for this library is  
CGCGTATACC. This library was created for the program, Gene  
Discovery in the Visual System, supported by National Eye  
Institute (NEI)."



/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-strand cDNA was digested with Not I and cloned into the Not I and EcoRV sites of the pCMVSPORT 6 vector. Library was not normalized."

ORIGIN

Alignment Scores:

Pred. No.:	6.56e-05	Length:	1201
Score:	98.00	Matches:	19
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	98.99%	Indels:	0
DB:	13	Gaps:	0

US-10-799-747-116 (1-20) x BX441923 (1-1201)

QY 1 MetAlaLaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19

Db 146 ATGGCAGCCCAATCAGCTTGAGTTTCTTCCTCCTGGACACCTTATGCTCTGAATCA 202

RESULT 6

AA654972

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

CONTACT: Robert Strausberg, Ph.D.  
Email: cgapbs-remail.nih.gov  
Tissue Procurement: W. Marston Linehan, M.D., Rodrigo Chuaqui, M.D., Michael Emmert-Buck, M.D., Ph.D.  
cDNA Library Preparation: David B. Krizman, Ph.D.  
cDNA Library Arrayed by: Genome Systems Inc., Greg Lennon, Ph.D.  
DNA Sequencing by: Washington University Genome Sequencing Center  
Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: www-bio.llnl.gov/bbrp/image/image.html  
Seq primer: -40ml3 fwd. ET from Amersham.

FEATURES

source

1..352

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="IMAGE:1203016"

/sex="Male"

/dev\_stage="45 years old"

/lab\_host="DH10B"

/clone\_lib="NCI CGAP Pr3"

/note="Vector: pAMP10; Site 1: NotI; Site 2: EcoRI; 1st strand cDNA was primed with oligo(dT)17 on 50 ng of DNase-treated, total cellular RNA obtained from 5,000-10,000 microdissected cells histologically determined to be fully malignant prostate cancer cells. Double-stranded cDNA was ligated to EcoRI adaptors, 5 cycles of PCR applied to the cDNA with an adaptor-specific primer, and the resulting PCR product subcloned into pAMP10 by the UDG-cloning method (Life Technologies). Average insert size is 600 bp. NOTE: Not directionally cloned. This library was constructed by David Krizman."

ORIGIN

Alignment Scores:

Pred. No.:	57.7	Length:	352
Score:	56.00	Matches:	9
Percent Similarity:	80.00%	Conservative:	3
Best Local Similarity:	60.00%	Mismatches:	3
Query Match:	56.57%	Indels:	0
DB:	9	Gaps:	0

US-10-799-747-116 (1-20) x AA654972 (1-352)

QY 4 HisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLys 18

Db 11 CACAGCAATATCATTCATTATTGGTGGCCACAGGCCATTAG 55

RESULT 7

CE789116/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

PUBMED

COMMENT

CONTACT: Kirkness EP  
The Institute for Genomic Research  
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive, Rockville, MD 20850, USA  
Tel: 301-838-0200  
Fax: 301-838-0208  
Email: ekirkness@tigr.org  
Class: shotgun.

FEATURES

source

1..642

/organism="Canis familiaris"

/mol\_type="genomic DNA"

/strain="Standard Poodle"

/db\_xref="taxon:9615"

/clone\_lib="Dog Library"

/note="Site 1: BstXI; Libraries were prepared from peripheral blood"

ORIGIN

Alignment Scores:

Pred. No.:	180	Length:	642
Score:	55.00	Matches:	10
Percent Similarity:	84.62%	Conservative:	1
Best Local Similarity:	76.92%	Mismatches:	2
Query Match:	55.56%	Indels:	0
DB:	29	Gaps:	0

US-10-799-747-116 (1-20) x CE789116 (1-642)

QY 7 LeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19

Db 208 TTATCATTCCTCCCTTGGTCCATTTTATTATANAAGC 170

RESULT 8

AZ831423

LOCUS

DEFINITION

ACCESSION

ORIGIN

```

VERSION      AZ831423.1  GI:13001331
KEYWORDS
SOURCE       Mus musculus (house mouse)
ORGANISM     Mus musculus
REFERENCE    1 (bases 1 to 723)
AUTHORS      Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,
            Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T.,
            Reilly,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von
            Niederhausen,A. and Wright,D., Weiss,R.
TITLE        Mouse whole genome scaffolding with paired end reads from 10kb
            plasmid inserts
JOURNAL
COMMENT      Unpublished (2000)
            Contact: Robert B. Weiss
            University of Utah Genome Center
            University of Utah
            Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
            84112, USA
            Tel: 801 585 5606
            Fax: 801 585 7177
            Email: ddunn@genetics.utah.edu
            Insert Length: 10000 Std Error: 0.00
            Plate: 0111 row: H column: 12
            Seq primer: CGTTGTAAACGCGGCAGT
            Class: plasmid ends
            High quality sequence stop: 723.
FEATURES     source
             1..723
                /organism="Mus musculus"
                /mol_type="genomic DNA"
                /strain="C57BL/6J"
                /db_xref="taxon:10090"
                /clone="UUGC2M0111H12"
                /sex="Male"
                /lab_host="E. Coli strain XL10-Gold, T1-resistant, F-"
                /clone_lib="Mouse 10kb plasmid UUGC1M library"
                /notes="Vector: PWD42nv; Purified genomic DNA from M.
            Laboratory Mouse DNA Resource
            (http://www.jax.org/resources/documents/dnares/). The DNA
            was hydrodynamically sheared by repeated passage through a
            0.005 inch orifice at constant velocity. The sheared DNA
            was blunt end-repaired with T4 DNA polymerase and T4
            polynucleotide kinase. Adaptor oligonucleotides were
            ligated to the blunt ends in high molar excess. The
            adapted DNA was purified and size-selected for a 9.5 to
            10.5 kb range using preparative agarose gel
            electrophoresis. Vector DNA was prepared from a derivative
            of PWD42 (gil4732114|gb|AF129072.1), a copy-number
            inducible derivative of plasmid R1. The vector was ligated
            with adaptors complementary to the insert adaptors and
            purified. The sheared, adapted mouse DNA was annealed to
            adapted vector DNA, and transformed into
            chemically-competent E. coli XL10-Gold (Stratagene) cells
            and selected for ampicillin resistance."
ORIGIN
Alignment Scores:
Pred. No.:      210      Length:      723
Score:          55.00    Matches:      9
Percent Similarity: 66.67% Conservative: 3
Best Local Similarity: 50.00% Mismatches: 6
Query Match:    55.56% Indels:      0
DB:             28      Gaps:        0

US-10-799-747-116 (1-20) x AZ831423 (1-723)

QY      1 MetAlaHisSerValLeuSerPheLeuLeuTrrProTyrAlaLeuLys 18
      ||| :||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      121 ATGCTCTCTCACAGCCTTGCAATTACTTGTGTGGTACCTTATCCCTTAG 174

RESULT 9

```

```

TA226G11P/c
LOCUS
DEFINITION   T. brucei sheared genomic DNA clone 226g11, forward sequence,
            genomic survey sequence.
ACCESSION   AL480086
VERSION     AL480086.1  GI:11845965
KEYWORDS    GSS.
SOURCE      Trypanosoma brucei
ORGANISM    Trypanosoma brucei
REFERENCE    1 (bases 1 to 550)
AUTHORS      Hall,N., Bowman,S., Lennard,N.J., Doggett,J., Atkin,R.,
            Chillingworth,C., Ormond,D., Harris,B., El-Sayed,N., Hou,L.,
            Melville,S.E., Rajandream,M.A. and Barrell,B.G.
TITLE        Direct Submission
JOURNAL
COMMENT      Submitted (10-DEC-2000) Trypanosoma brucei genome sequencing
            project, Sanger Centre, The Wellcome Trust Genome Campus, Hinxton,
            Cambridge CB10 1SA, E-mail: barrell@sanger.ac.uk and
            nh@sanger.ac.uk
            Constructed at the Institute for Genomic Research (TIGR),
            Rockville, MD. Genomic DNA isolated from a cloned population of
            Trypanosoma brucei (TREU927/4 Gurat 10.1) was mechanically sheared
            to give a tight size distribution (
            4 kb). The v + i method used for the library construction is
            described in detail in Smith, H. and Venter, J.C. (Making small
            insert libraries for whole genome shotgun sequencing projects. In
            Genome Sequencing: A Practical Approach, eds. M. Vaudin and B.
            Barrell, Oxford University Press, 1999).
            Email: nh@sanger.ac.uk
            Details of T. brucei sequencing at the Sanger Centre are available
            at http://www.sanger.ac.uk/Projects/T_brucei/.
FEATURES     source
             1..550
                /organism="Trypanosoma brucei"
                /mol_type="genomic DNA"
                /strain="TREU927"
                /db_xref="taxon:5691"
                /clone="226g11"
ORIGIN
Alignment Scores:
Pred. No.:      212      Length:      550
Score:          54.00    Matches:      11
Percent Similarity: 73.33% Conservative: 0
Best Local Similarity: 73.33% Mismatches: 4
Query Match:    54.55% Indels:      0
DB:             29      Gaps:        0

US-10-799-747-116 (1-20) x TA226G11P (1-550)

QY      3 AlaHisSerValLeuSerPheLeuLeuTrrProTyrAlaLeu 17
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      356 GCGGGTGGTGTGCTTGTAGTTTTTTTGTGGACTCTTAGCGCGCTG 312

RESULT 10
AQ645123
LOCUS
DEFINITION   RPC193-EcoRI-2M16-TV RPC193-EcoRI Trypanosoma brucei genomic clone
            genomic survey sequence.
ACCESSION   AQ645123
VERSION     AQ645123.1  GI:5121833
KEYWORDS    GSS.
SOURCE      Trypanosoma brucei
ORGANISM    Trypanosoma brucei
REFERENCE    1 (bases 1 to 724)
AUTHORS      El-Sayed,N., Zhao,S., Zhao,H., Gill,S., Suh,E., Malek,J., Fujii,C.,
            Gerrard,C., Leech,V., de Jong,P., Ullu,E., Melville,S.,
            Donelson,J., Fraser,C. and Adams,M.
            Use of BAC end sequences from Trypanosoma brucei GUTat 10.1 RPCI-93
            Library for gene discovery and sequence-ready map construction

```

JOURNAL  
COMMENT

Unpublished (1999)  
Other GSSs: RPC193-EcoRI-2M16.TP  
Contact: Najib M. El-Sayed  
Department of Eukaryotic Genomics  
The Institute for Genomic Research  
9712 Medical Center Dr., Rockville, MD 20850, USA  
Tel: 301 838 0200  
Fax: 301 838 0208  
Email: nelsayed@tigr.org  
Clones and high density filters may be purchased from BACPAC  
Resources (http://bacpac.med.buffalo.edu). BAC end sequences search  
page: http://www.tigr.org/tdb/mbd/tbdb/.

Seq primer: T7

Class: BAC ends.

FEATURES  
source

Location/Qualifiers  
1..724  
/organism="Trypanosoma brucei"  
/mol\_type="genomic DNA"  
/strain="TREU927/4 GUTat 10.1"  
/db\_xref="taxon:5691"  
/clone="RPC193-EcoRI-2M16"  
/note="Vector: pBAC3.6; Site 1: Eco RI; Site 2: Eco RI;  
Constructed for The Institute for Genomic Research by  
Bohui Zhao in Pieter de Jong's laboratory (Roswell Park  
Cancer Institute, Buffalo, NY). Briefly, Trypanosoma  
brucei TREU927/4 GUTat 10.1 agarose embedded DNA was  
partially digested with a combination of Eco RI and Eco RI  
methylase (RPC193-EcoRI segment) or Dpn II (RPC193-DpnII  
segment). High molecular weight fragments were ligated in  
pBAC3.6 vector digested with Eco RI or Bam HI,  
respectively. The average insert size is 141 Kb. Total  
coverage (both segments): > 90 X the haploid  
non-minichromosomal genome."

## ORIGIN

Alignment Scores:  
Pred. No.: 302 Length: 724  
Score: 54.00 Matches: 11  
Percent Similarity: 73.33% Conservative: 0  
Best Local Similarity: 73.33% Mismatches: 4  
Query Match: 54.55% Indels: 0  
DB: 28 Gaps: 0

US-10-799-747-116 (1-20) x AQ645123 (1-724)

Qy 3 AlahisSerValLeuSerPheLeuLeuTrpThrProTyrAlaLeu 17

Db 329 GCGGGTGTGCTAGTTTGTGGTCTCTTTAGGCTG 373

## RESULT 11

CA967853 816 bp mRNA linear EST 03-JAN-2003  
DEFINITION CcLX04a13k03f1 Carp mixed tissue library 1 Cyprinus carpio cDNA  
clone 13k03 5', mRNA sequence.

ACCESSION CA967853

VERSION CA967853.1 GI:27494410

KEYWORDS EST.

SOURCE Cyprinus carpio (common carp)

## ORGANISM

Cyprinus carpio  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Actinopterygii; Neopterygii; Teleostei; Ostariophysi;  
Cypriniformes; Cyprinidae; Cyprinus.

1 (bases 1 to 816)

REFERENCE Gracey,A.Y., Fraser,E., Li,W. and Cossins,A.R.

AUTHORS Microarray and EST analysis of the carp (Cyprinus carpio)

TITLE transcriptome during environmental stress

JOURNAL Unpublished (2003)

COMMENT Contact: Andrew R. Cossins

Laboratory for Environmental Gene Regulation

University of Liverpool

School of Biological Sciences, The Biosciences Building, Crown

Street, Liverpool, United Kingdom, L69 7ZB

Tel: +44 (0)151-795-4510  
Fax: +44 (0)151-795-4431  
Email: cossins@liverpool.ac.uk  
Vector has been trimmed from this EST.  
Plate: 13 row: k column: 03  
Seq primer: Triplex 5' LD (5'-CTCGGAGCGCGCCATTGTGTGGT-3')  
High quality sequence start: 33  
High quality sequence stop: 494.

FEATURES  
source

Location/Qualifiers  
1..816  
/organism="Cyprinus carpio"  
/mol\_type="mRNA"  
/db\_xref="taxon:7962"  
/clone="13k03"  
/sex="Male & female"  
/tissue\_type="Skeletal white muscle, cardiac muscle,  
kidney, brain, gill, intestinal mucosa"  
/dev\_stage="Adult"  
/lab\_host="E.coli Electromax DH10B"  
/clone\_lib="Carp mixed tissue library 1"  
/note="Vector: pTriplex2; Site 1: SfiI GGCCATTACGGCC;  
Site 2: SfiI GGCGGCTCGGC; Normalized cDNA library  
prepared from mixed tissues of warm, cold and hypoxia  
challenged animals"

## ORIGIN

Alignment Scores:  
Pred. No.: 353 Length: 816  
Score: 54.00 Matches: 10  
Percent Similarity: 80.00% Conservative: 2  
Best Local Similarity: 66.67% Mismatches: 3  
Query Match: 54.55% Indels: 0  
DB: 14 Gaps: 0

US-10-799-747-116 (1-20) x CA967853 (1-816)

Qy 5 SerValLeuSerPheLeuLeuTrpThrProTyrAlaLeuLysSer 19

Db 679 TCTGCTCTCTTTCTTACTTTGGTTCCTTCCTCTTCTTC 723

## RESULT 12

AA766276/c

LOCUS

DEFINITION

AA766276

oa29c04.s1 NCI CGAP GCB1 Homo sapiens cDNA

similar to contains element OPR repetitive element ;, mRNA

sequence.

ACCESSION AA766276

VERSION AA766276.1 GI:2817514

KEYWORDS EST.

SOURCE Homo sapiens (human)

## ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 394)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.

Email: ccapbs-remail.nih.gov

Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,

Ph.D., Gerald Marti, M.D.

cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 666 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 357.

Location/Qualifiers

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source
1. 394
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/db_xref="taxon:9606"
/clone="IMAGE:1306374"
/tissue_type="germinal center B cell"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_GCB1"
/notes="vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from human tonsillar cells enriched for
germinal center B cells by flow sorting (CD20+, Igu-),
provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
(NCI) and Dr. Gerald Marti (CSER). cDNA synthesis was
primed with a Not I - oligo (dT) primer
[5'-TGTTACCAATCTCAAGTGGAGCGCGCTCATTTTTTTTTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN
Alignment Scores:
Pred. No.: 199 Length: 394
Score: 53.00 Matches: 9
Percent Similarity: 80.00% Conservative: 3
Best Local Similarity: 60.00% Mismatches: 3
Query Match: 53.54% Indels: 0
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AA804444 (1-394)

Qy 2 AlaAlaHisSerValLeuSerPheLeuLeuTrpThrProTyrAla 16
|||||
Db 166 GCTGCACACTCTCTCTTTTATTCTCTGGCCCAATTTCC 122

RESULT 13
AA804444/c
LOCUS
DEFINITION n238c04.s1 NCI CGAP GCB1 Homo sapiens cDNA clone IMAGE:1290054 3'
similar to contains element QFR repetitive element ;, mRNA
sequence.
AA804444 429 bp mRNA linear EST 18-FEB-1998
VERSION AA804444.1 GI:2875957
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 429)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT Contact: Robert Strausberg, Ph.D.
Email: cgaps-remail.nih.gov
Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
Ph.D., Gerald Marti, M.D.
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
Bonaldo, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 649 Std Error: 0.00
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 249.
Location/Qualifiers
1. 429
/organism="Homo sapiens"
/mol_type="mRNA"

/db_xref="taxon:9606"
/clone="IMAGE:1290054"
/tissue_type="germinal center B cell"
/lab_host="DH10B"
/clone_lib="NCI_CGAP_GCB1"
/notes="vector: pT7T3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from human tonsillar cells enriched for
germinal center B cells by flow sorting (CD20+, Igu-),
provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
(NCI) and Dr. Gerald Marti (CSER). cDNA synthesis was
primed with a Not I - oligo (dT) primer
[5'-TGTTACCAATCTCAAGTGGAGCGCGCTCATTTTTTTTTTTT-3'
]. Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization, and was
constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN
Alignment Scores:
Pred. No.: 222 Length: 429
Score: 53.00 Matches: 9
Percent Similarity: 80.00% Conservative: 3
Best Local Similarity: 60.00% Mismatches: 3
Query Match: 53.54% Indels: 0
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AA804444 (1-429)

Qy 2 AlaAlaHisSerValLeuSerPheLeuLeuTrpThrProTyrAla 16
|||||
Db 166 GCTGCACACTCTCTCTTTTATTCTCTGGCCCAATTTCC 122

RESULT 14
B1140807/c
LOCUS
DEFINITION B1140807 600 bp mRNA linear EST 03-JUL-2001
mRNA sequence.
B1140807
VERSION B1140807.1 GI:14593250
KEYWORDS EST.
SOURCE Sorghum bicolor (sorghum)
ORGANISM Sorghum bicolor
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; PACAD
clade; Panicoideae; Andropogoneae; Sorghum.
1 (bases 1 to 600)
Klein,R.R., Cordonnier-Pratt,M.-M., Gingle,A., Sudman,M. and
Pratt,L.H.
An EST database from Sorghum: developing preanthesis pannicles
Unpublished (2001)
Contact: Cordonnier-Pratt MM
Laboratory for Genomics and Bioinformatics
The University of Georgia, Department of Plant Biology
Plant Sciences Building, Rm. 2502, Athens, GA 30602-7271, USA
Tel: 706 542 1860
Fax: 706 583 0210
Email: mmpratt@uga.edu
Sequences have been trimmed to exclude PolyA, vector and regions
below Phred quality 16. The threshold for high quality sequence is
20. Three-prime sequences, which are obtained with PolyTMix or T7
sequencing primer, are presented as the reverse complement.
Seq primer: JEN REV
High quality sequence stop: 538
POLYA=No.
Location/Qualifiers
1. 600
/organism="Sorghum bicolor"
/mol_type="mRNA"
/cultivar="BTx623"
/db_xref="taxon:4558"
/clone_lib="Immature pannicle 1 (IPI)"
FEATURES
source

```





GenCore version 5.1.6  
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 02:54:41 ; Search time 1704 Seconds  
(without alignments)

508.721 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 20

Sequence: 1 MAHSVLSFLWTPYALKSX 20

Scoring table: OLIGO  
Xgapop 60.0 , Xgapext 60.0  
Ygapop 60.0 , Ygapext 60.0  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 3470272 seqs, 21671516995 residues

Word size: 1

Total number of hits satisfying chosen parameters: 6933152

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Command line parameters: -DEV=xlpl  
-MODEL=frame\_p2n.model  
-Q=/cgn2\_1/USPTO\_spool\_p/US10799747/runat\_19072004\_161442\_21981/app\_query.fasta\_1.199  
-DB=GenEmbl -QFMT=fastap -SURFIX=olig.rge -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0  
-UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi -LIST=45  
-DOCALIGN=200 -THR SCORE=quality -THR MIN=1 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc  
-NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10799747@cgn1\_1.5600 @runat\_19072004\_161442\_21981 -NCFU=6 -ICPU=3  
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6  
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database : GenEmbl.\*  
1: gb.ba.\*  
2: gb.htg.\*  
3: gb.in.\*  
4: gb.om.\*  
5: gb.ov.\*  
6: gb.pat.\*  
7: gb.ph.\*  
8: gb.pl.\*  
9: gb.pr.\*  
10: gb.ro.\*  
11: gb.sts.\*  
12: gb.sy.\*  
13: gb.un.\*  
14: gb.vi.\*  
15: em.ba.\*  
16: em.fun.\*  
17: em.hum.\*  
18: em.in.\*  
19: em.mu.\*  
20: em.om.\*  
21: em.or.\*  
22: em.ov.\*  
23: em.pat.\*  
24: em.ph.\*  
25: em.pl.\*  
26: em.ro.\*  
27: em.sts.\*  
28: em.un.\*

29: em.vi.\*  
30: em.htg.hum.\*  
31: em.htg.inv.\*  
32: em.htg.other.\*  
33: em.htg.mus.\*  
34: em.htg.pln.\*  
35: em.htg.fod.\*  
36: em.htg.mam.\*  
37: em.htg.vrt.\*  
38: em.sy.\*  
39: em.htgo.hum.\*  
40: em.htgo.mus.\*  
41: em.htgo.other.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	19	95.0	1434	6	BD078422	BD078422 101 human
2	19	95.0	3452	9	BC033650	BC033650 Homo sapi
3	19	95.0	3522	9	HSM804706	AL833393 Homo sapi
4	19	95.0	175081	9	AC021151	AC021151 Homo sapi
5	19	95.0	281662	2	AC117374	AC117374 Homo sapi
6	13	65.0	2020	9	HSM800227	AL049442 Homo sapi
7	9	45.0	153464	2	AC123991	AC123991 Mus muscu
8	9	45.0	167013	9	HSDJ828H9	AL121757 Human DNA
9	9	45.0	205725	10	AC122809	AC122809 Mus muscu
10	9	45.0	231768	2	AC110315	AC110315 Rattus no
11	9	45.0	254839	2	AC103176	AC103176 Rattus no
12	9	45.0	267352	2	AC098074	AC098074 Rattus no
13	9	45.0	306660	2	AC098226	AC098226 Rattus no
14	8	40.0	339	11	G25673	G25673 human STS E
15	8	40.0	356	11	G27758	G27758 human STS S
16	8	40.0	691	3	AK112944	AK112944 Clona int
17	8	40.0	1503	6	AX654294	AX654294 Sequence
18	8	40.0	1524	9	AB070060	AB070060 Macaca fa
19	8	40.0	1751	9	AK130941	AK130941 Homo sapi
20	8	40.0	1791	8	AK106862	AK106862 Oryza sat
21	8	40.0	1806	10	MMY15799	Y15799 Mus musculu
22	8	40.0	1858	10	MMY15797	Y15797 Mus musculu
23	8	40.0	1916	10	MMY15800	Y15800 Mus musculu
24	8	40.0	1952	10	AF040749	AF040749 Mus muscu
25	8	40.0	1975	8	AY090039	AY090039 Nicotiana
26	8	40.0	1997	10	AF040747	AF040747 Mus muscu
27	8	40.0	1999	10	AF040748	AF040748 Mus muscu
28	8	40.0	2841	10	MMY15798	Y15798 Mus musculu
29	8	40.0	3200	8	AK119280	AK119280 Oryza sat
30	8	40.0	3628	8	AF238476	AF238476 Oryza sat
31	8	40.0	14782	9	HSMLADPB	X02228 Human HLA-D
32	8	40.0	37201	9	AP001215	AP001215 Homo sapi
33	8	40.0	52058	2	AC100335	AC100335 Mus muscu
34	8	40.0	56083	2	AC141162	AC141162 Rattus no
35	8	40.0	60504	2	AC090716	AC090716 Homo sapi
36	8	40.0	62263	2	AC100239	AC100239 Mus muscu
37	8	40.0	62263	2	AC100239	AL833790 Mouse DNA
38	8	40.0	63139	10	AL833790	AL833790 Mouse DNA
39	8	40.0	64380	9	BX120009	BX120009 Human DNA
40	8	40.0	65209	2	AC101355	AC101355 Mus muscu
41	8	40.0	67358	2	AC108426	AC108426 Mus muscu
42	8	40.0	68790	9	AF017732	AF017732 Homo sapi
43	8	40.0	74658	8	OS243961	AJ243961 Oryza sat
44	8	40.0	91585	10	AL606661	AL606661 Mouse DNA
45	8	40.0	93842	9	AL845446	AL845446 Human DNA

ALIGNMENTS

```

BD0784422
LOCUS      BD078422      1434 bp      DNA      linear      PAT 27-AUG-2002
DEFINITION 101 human secretory proteins.
ACCESSION  BD078422
VERSION    BD078422.1 GI:22624025
KEYWORDS   JP 2001519156-A/11.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 1434)
AUTHORS   Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
            Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
            Altschul,S.F., Zeeberg,B., Buetow,K.H., Stoenen,C.F., Bhat,N.K.,
            Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
            Diachenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
            Schapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
            Scheetz,T.E., Brownstein,M.J., Utsdin,T.B., Toshiyuki,S.,
            Carninci,P., Prange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
            Abramson,R.D., Mullahy,S.J., Bosak,S.A., McManus,P.J.,
            McKernan,K.O., Malek,J.A., Gunaratne,P.H., Richards,S.,
            Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W.,
            Villalón,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
            Fahey,J., Helton,E., Kettman,M., Madan,A., Young,A.C., Shvachenko,Y.,
            Sanchez,A., Whiting,M., Madan,A., Young,A.C., Shvachenko,Y.,
            Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
            Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
            Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalius,D.E.,
            Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
            Generation and initial analysis of more than 15,000 full-length
            human and mouse cDNA sequences
            Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
2238257
JOURNAL    MEDLINE
PUBMED     12477932
REFERENCE  2 (bases 1 to 3452)
AUTHORS   Strausberg,R.
TITLE     Direct Submission
JOURNAL
REMARK
COMMENT    NIH-MGC Project URL: http://mgc.nci.nih.gov
            Contact: MGC help desk
            Email: cgabs@mail.nih.gov
            Tissue Procurement: ATCC
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: National Institutes of Health Intramural
            Sequencing Center (NISC),
            Gaithersburg, Maryland;
            Web site: http://www.nisc.nih.gov/
            Contact: nisc.mgc@nih.gov
            Akhter,N., Ayele,K., Beckstrom-Sternberg,S.M., Benjamin,B.,
            Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S.,
            Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P.,
            Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Laric,P., Legaapi,R.,
            Maduro,Q.L., Masiello,C., Maskeri,B., Mastrian,S.D., McCloskey,J.C.,
            McDowell,J., Pearson,R., Stantripop,S., Thomas,P.J., Touchman,J.W.,
            Tsurgeon,C., Vogt,J.D., Walker,M.A., Wetherby,K.D., Wiggins,L.,
            Young,A., Zhang,L.-H. and Green,E.D.
            Clone distribution: MGC clone distribution information can be found
            through the I.M.A.G.E. Consortium/LLNL at: http://image.llnl.gov
            Series: IRAK Plate: 69 Row: n Column: 19
            This clone was selected for full length sequencing because it
            passed the following selection criteria: matched mRNA gi: 14249445.
FEATURES   source
            1..3452
            /location/Qualifiers
            /organism="Homo sapiens"
            /mol_type="mRNA"
            /db_xref="taxon:9606"
            /clone="MGC:44889 IMAGE:5574637"
            /tissue type="Duodenum, adenocarcinoma"
            /clone_lib="NIH MGC_88"
            /lab host="DH10B"
            /note="vector: pCMV-SPORT6"
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            /db_xref="LocusID:84869"
            159..872
            CDS

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/codon_start=1
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ALAKVARKIRVNVVAPGVFHTMTKDLKEBLKKNIPLGRFGETIEVAHAVFELLE
SPYITGVLVLDGGLQLIL"
misc_feature
159..857
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family contains a wide variety of dehydrogenases"
/db_xref="CDD:pfam00106"

ORIGIN
Alignment Scores:
Pred. No.: 6.82e-10 Length: 3452
Score: 19.00 Matches: 19
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 95.00% Indels: 0
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x BC033650 (1-3452)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19
|||||
DB 2542 ATGGCAGCCCAATTCAGTCTTGAGTTTCTTCTCTGGACACCTTATGCTCTGAAATCA 2598

RESULT 3
HSM804706
LOCUS Homo sapiens mRNA; cDNA DKFp762K109 (from clone DKFp762K109).
ACCESSION AL833393
VERSION AL833393.1 GI:21734029
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM
REFERENCE
AUTHORS Kyung,Y. and Abbott,A.
TITLE The sequence of Homo sapiens BAC clone RP11-483A20
JOURNAL Unpublished (2001)
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (14-JUN-2000) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63106, USA
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (09-AUG-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63106, USA
REFERENCE
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (09-JAN-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Aug 9, 2001 this sequence version replaced gi:13877272.
COMMENT
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@wustl.wustl.edu
----- Summary Statistics
-----
Center project name: H_NH0483A20
-----

NOTICE: This sequence may not represent the entire insert of this
clone. It may be shorter because we only sequence overlapping
clone sections once, or longer because we provide a small overlap
between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate
chemistry, or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by sequence
from more than one subclone; and the assembly was confirmed by
restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D.
```

McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

## SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Woon, P.Y., Zhao, B., Frengen, E., Tateno, M., Catanese, J.J. and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

## NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the right is RP11-36G9. Actual start of this clone is at base position 1 of RP11-483A20; actual end is at base position 175081 of RP11-483A20.

## FEATURES

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3270..3453	
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3434..3464	
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3455..4161	
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7949..7956	
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10495..10665	
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10731..11051	
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## Alignment Scores:

Pred. No.:	2,42e-08	Length:	175081
Score:	19.00	Matches:	19
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	95.00%	Indels:	0

DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AC021151 (1-175081)

QY 1 MetAlaLaHisSerValLeuSerPheLeuLeuTrpThrProTyrAlaLeuLysSer 19  
|||||

Db 57641 ATGCACGCCATTCAGTCTTGAGTTTCTCTCGGACACCTTAAGTCTCTGAATCA 57585

RESULT 5  
AC117374/c  
LOCUS  
DEFINITION Homo sapiens chromosome 12 clone RP11-19D19, \*\*\* SEQUENCING IN  
ACCESSION AC117374 AC011694  
VERSION AC117374.1 GI:20127352  
KEYWORDS HTG; HTGS PHASE1.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens

REFERENCE  
AUTHORS  
1 (bases 1 to 281662)  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-Osman,F.R., Allen,C.,  
Alsbrooks,S.L., Amarantunga,H.C., Are,J.R., Ayele,M., Banks,T.,  
Barbaria,J., Benton,J., Bimage,K., Blankenburg,K., Bonnin,D.,  
Bouck,J., Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P.,  
Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,  
Carroll,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,  
Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C.,  
Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,  
Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,  
Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H.,  
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Rives,M., Rojas,A., Rojubokan,I., Rolfe,M., Ruiz,S., Savery,G.,  
Scherer,S., Scott,G., Shen,H., Shooshtari,N., Sisson,I.,  
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Tansley,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,  
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Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,  
Williams,G., Williamson,A., Wleczyk,R., Wooden,S., Worley,K.,  
Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,  
Weinstock,G. and Gibbs,R.

Direct Submission  
Unpublished  
2 (bases 1 to 281662)  
Worley,K.C.  
Direct Submission  
Submitted (10-APR-2002) Human Genome Sequencing Center, Department  
of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
3 (bases 1 to 281662)  
Worley,K.C.  
Direct Submission  
Submitted (02-MAY-2002) Human Genome Sequencing Center, Department

COMMENT

of Molecular and Human Genetics, Baylor College of Medicine, One  
Baylor Plaza, Houston, TX 77030, USA  
On Apr 28, 2002 this sequence version replaced gi:10045385.

Center: Baylor College of Medicine  
Center code: BCM  
Web site: <http://www.hgsc.bcm.tmc.edu/>  
Contact: hgsc-help@bcm.tmc.edu  
Project Information  
Center project name: HCNB  
Center clone name: RP11-19D19  
Summary Statistics  
Sequencing vector: Plasmid;  
Chemistry: Dye-terminator Big Dye; 100% of reads  
Assembly program: Phrap; version 0.990329  
Consensus quality: 273063 bases at least Q40  
Consensus quality: 294336 bases at least Q30  
Consensus quality: 306942 bases at least Q20

\* NOTE: Estimated insert size may differ from sequence length  
(see [http://www.hgsc.bcm.tmc.edu/docs/Genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)).  
\* NOTE: This is a 'working draft' sequence. It currently  
\* consists of 48 contigs. The true order of the pieces  
\* is not known and their order in this sequence record is  
\* arbitrary. Gaps between the contigs are represented as  
\* runs of N, but the exact sizes of the gaps are unknown.  
\* This record will be updated with the finished sequence  
\* as soon as it is available and the accession number will  
\* be preserved.

1 2464: contig of 2464 bp in length  
\* 2465: gap of unknown length  
\* 2565: contig of 2039 bp in length  
\* 4603: gap of unknown length  
\* 4704: contig of 2410 bp in length  
\* 7113: gap of unknown length  
\* 7213: gap of unknown length  
\* 9517: contig of 2304 bp in length  
\* 9518: gap of unknown length  
\* 9618: contig of 2490 bp in length  
\* 12108: gap of unknown length  
\* 12208: contig of 2807 bp in length  
\* 15015: gap of unknown length  
\* 15115: contig of 2717 bp in length  
\* 17832: gap of unknown length  
\* 17931: contig of 2549 bp in length  
\* 20480: gap of unknown length  
\* 20581: contig of 2120 bp in length  
\* 22701: gap of unknown length  
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\* 25010: contig of 2304 bp in length  
\* 27314: gap of unknown length  
\* 27414: contig of 2385 bp in length  
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\* 78107 78107: contig of 4562 bp in length  
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\* 116222 116321: gap of unknown length  
\* 116322 113798: contig of 3477 bp in length  
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\* 119899 124870: contig of 4972 bp in length  
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\* 130791 130890: gap of unknown length  
\* 130891 134639: contig of 3749 bp in length  
\* 134640 134739: gap of unknown length  
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\* 174993 184400: contig of 9408 bp in length  
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FEATURES

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ORIGIN

Alignment Scores:  
Pred. No.: 3.73e-08 Length: 281662  
Score: 19.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 95.00% Indels: 0  
DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC117374 (1-281662)

Qy 1 MetAlaLaHisSerValLeuSerPheLeuLeuTrpThrProTyAlaLeuIysSer 19

Db 145718 ATGGCAGCCCATTCAGTCTTGAGTTTCTTCTCTGGACCATATGCTCTGAAATCA 145662

RESULT 6

HSM800227 2020 bp mRNA linear PRI 18-FEB-2000  
LOCUS Homo sapiens mRNA; cDNA DKFZp586N1720 (from clone DKFZp586N1720).  
DEFINITION  
ACCESSION AL049442  
VERSION AL049442.1 GI:4500222  
KEYWORDS  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE  
1 (bases 1 to 2020)  
AUTHORS Wambutt,R., Heubner,D., Mewes,H.W., Gassenhuber,J. and Wiemann,S.  
TITLE Direct Submission  
JOURNAL Submitted (10-MAR-1999) MIPS, Am Klopferspitz 18a, D-82152  
COMMENT Martinsried, GERMANY  
Clone from S. Wiemann, Molecular Genome Analysis, German Cancer  
Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;  
sequenced by AGOWA (Berlin/Germany) within the cDNA sequencing  
consortium of the German Genome Project.  
This clone (DKFZp586N1720) is available at the RZPD in Berlin.  
Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059  
Berlin-Charlottenburg, GERMANY; Email: clone@zpd.de Further  
information about the clone and the sequencing project is available  
at http://www.mips.biochem.mpg.de/proj/cDNA/.

FEATURES

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ORIGIN

Alignment Scores:  
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Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 65.00% Indels: 0  
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x HSM800227 (1-2020)

Qy 1 MetAlaLaHisSerValLeuSerPheLeuLeuTrpThr 13

Db 1115 ATGGCAGCCCATTCAGTCTTGAGTTTCTTCTCTGGACA 1153

RESULT 7

AC123991/c 153464 bp DNA linear HTG 17-MAR-2003  
LOCUS Mus musculus clone RP24-137I10, WORKING DRAFT SEQUENCE, 3 unordered  
DEFINITION  
ACCESSION AC123991  
VERSION AC123991.3 GI:28975888  
KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT.  
SOURCE Mus musculus (house mouse)  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE  
1 (bases 1 to 153464)  
AUTHORS Birren,B., Nusbaum,C. and Lander,E.  
TITLE Mus musculus, clone RP24-137I10  
JOURNAL Unpublished  
REFERENCE 2 (bases 1 to 153464)  
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,

Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L.,  
 Boukhgalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,  
 Chazaro, B., Choepel, Y., Collangelo, M., Collins, S., Collymore, A.,  
 Cook, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S.,  
 Fero, S., Ferreira, P., Fitzgerald, M., FitzHugh, W., Gage, D.,  
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 Grand-Pierre, N., Hagos, B., Horton, L., Hulme, W., Iliev, I.,  
 Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., LaRoque, K.,  
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 Liu, G., MacLean, C., Macdonald, P., Major, J., Marquis, N.,  
 Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrim, J.,  
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 O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N.,  
 Pollar, V., Raymond, C., Retta, R., Rieback, M., Riley, R., Rise, C.,  
 Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S.,  
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 Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J.,  
 Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J.,  
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Direct Submission  
 Submitted (06-JUN-2002) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 3 (bases 1 to 153464)

Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,  
 Anderson, S., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T.,  
 Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Choepel, Y.,  
 Collymore, A., Cooke, P., Cooke, P., Corum, B., DeArellano, K.,  
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 Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M.,  
 Meldrim, J., Meneus, L., Mihova, T., Menga, V., Murphy, T., Naylor, J.,  
 Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P.,  
 O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N.,  
 Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P.,  
 Roman, J., Schauer, S., Schupback, R., Seaman, S., Severy, P., Smith, C.,  
 Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M.,  
 Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M.,  
 Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X.,  
 Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission  
 Submitted (17-MAR-2003) Whitehead Institute/MIT Center for Genome  
 Research, 320 Charles Street, Cambridge, MA 02141, USA  
 On Mar 17, 2003 this sequence version replaced gi:28201678.  
 All repeats were identified using RepeatMasker:  
 Smit, A.F.A. & Green, P. (1996-1997)  
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center  
 Center: Whitehead Institute/ MIT Center for Genome Research  
 Center code: WITR  
 Web site: <http://www-seq.wi.mit.edu>  
 Contact: [sequence\\_submissions@genome.wi.mit.edu](mailto:sequence_submissions@genome.wi.mit.edu)

----- Project Information  
 Center project name: L26544  
 Center clone name: 137.1.10  
 ----- Summary Statistics

Sequencing vector: Plasmid; n/a; 100% of reads  
 Chemistry: Dye-terminator Big Dye; 100% of reads  
 Assembly program: Phrap; version 0.960731

Consensus quality: 152581 bases at least Q40  
 Consensus quality: 152929 bases at least Q30  
 Consensus quality: 153074 bases at least Q20

Insert size: 151000; agarose-fp  
 Insert size: 153284; sum-of-contigs  
 Quality coverage: 11.8 in Q20 bases; agarose-fp  
 Quality coverage: 11.7 in Q20 bases; sum-of-contigs

-----  
 \* NOTE: This is a 'working draft' sequence. It currently

\* consists of 3 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

\* 1 132252: contig of 132252 bp in length  
 \* 132253 132352: gap of 100 bp  
 \* 132353 134109: contig of 1757 bp in length  
 \* 134110 134209: gap of 100 bp  
 \* 134210 153464: contig of 19255 bp in length.

## FEATURES

## source

1. 153464  
 /location/Qualifiers  
 1. 153464  
 /organism="Mus musculus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:10090"  
 /clone="RP24-137110"  
 /clone\_lib="RPCI-24 Male Mouse BAC"  
 1. 132252  
 /note="assembly\_fragment  
 clone\_end:SP6  
 vector\_side:left"  
 132353. 134109  
 /note="assembly\_fragment"  
 134210. 153464  
 /note="assembly\_fragment  
 clone\_end:T7  
 vector\_side:right"

## misc\_feature

1. 153464  
 /note="assembly\_fragment  
 clone\_end:SP6  
 vector\_side:left"  
 132353. 134109  
 /note="assembly\_fragment"  
 134210. 153464  
 /note="assembly\_fragment  
 clone\_end:T7  
 vector\_side:right"

## misc\_feature

1. 153464  
 /note="assembly\_fragment  
 clone\_end:SP6  
 vector\_side:left"  
 132353. 134109  
 /note="assembly\_fragment"  
 134210. 153464  
 /note="assembly\_fragment  
 clone\_end:T7  
 vector\_side:right"

## misc\_feature

1. 153464  
 /note="assembly\_fragment  
 clone\_end:SP6  
 vector\_side:left"  
 132353. 134109  
 /note="assembly\_fragment"  
 134210. 153464  
 /note="assembly\_fragment  
 clone\_end:T7  
 vector\_side:right"

## ORIGIN

## Alignment Scores:

Pred. No.: 326 Length: 153464  
 Score: 9.00 Matches: 9  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 45.00% Indels: 0  
 DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC123991 (1-153464)

Qy 3 ALAHISerValLeuSerPheLeuLeu 11

|||||  
 Db 102764 GCCACTCAGTACTGAGCTTTTATTG 102738

## RESULT 8

## HSDJ828H9/c

LOCUS HSDJ828H9 167013 bp DNA linear PRI 06-MAR-2001  
 DEFINITION Human DNA sequence from clone RP5-828H9 on chromosome 20. Contains  
 two putative novel genes, STSs, GSSs and CpG islands, complete  
 sequence.

## ACCESSION

## AL121757

## VERSION

## AL121757.7 GI:6114793

## KEYWORDS

## HTG; CpG island.

## SOURCE

## Homo sapiens

## ORGANISM

## Homo sapiens

## Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

## REFERENCE

## 1 (bases 1 to 167013)

## AUTHORS

## Blakey, S.

## TITLE

## Direct Submission

## JOURNAL

Submitted (28-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,  
 CB10 1SA, UK. E-mail enquiries: [humquery@sanger.ac.uk](mailto:humquery@sanger.ac.uk)  
 Requests: [clonerequest@sanger.ac.uk](mailto:clonerequest@sanger.ac.uk)

## COMMENT

## On Oct 25, 1999 this sequence version replaced gi:6102834.

## During sequence assembly data is compared from overlapping clones.

## Where differences are found these are annotated as variations

## together with a note of the overlapping clone name. Note that the

## variation annotation may not be found in the sequence submission

## corresponding to the overlapping clone, as we submit sequences with

## only a small overlap as described above.

## The following abbreviations are used to associate primary accession

## numbers given in the feature table with their source databases:

Em., EMBL; Sw., SWISSPROT; Tr., TREMBL; Wp., WORMPEP; Information on the WORMPEP database can be found at [http://www.sanger.ac.uk/Projects/C\\_elegans/wormpep](http://www.sanger.ac.uk/Projects/C_elegans/wormpep) This sequence was generated from part of bacterial clone contigs of human chromosome 20, constructed by the Sanger Centre Chromosome 20 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr20>

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. RP5-828H9 is from the library RPI-5 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

VECTOR: pCYPAC2

IMPORTANT: This sequence is not the entire insert of clone RP5-828H9 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The left end of clone RP4-680N4 is at 166918 in this sequence. The true left end of clone RP5-1022P6 is at 99 in this sequence. The true right end of clone RP4-766D4 is at 46185 in this sequence.

Location/Qualifiers

1..167013

/organism="Homo sapiens"

/mol\_type="genomic DNA"

/db\_xref="taxon:9606"

/chromosome="20"

/clone="RP5-828H9"

/clone\_lib="RPI-5"

785..1214

/note="HERV17 repeat: matches 6052..6478 of consensus"

1271..1441

/note="match: GSS: Em:AQ780632"

1336..1566

/note="MER66-internal repeat: matches 4624..4847 of consensus"

1446..1607

/note="match: GSS: Em:AQ780632"

1798..1923

/note="MER51-internal repeat: matches 7690..7816 of consensus"

2182..2263

/note="MER51B repeat: matches 190..523 of consensus"

2375..2451

/note="AluS repeat: matches 1..67 of consensus"

2455..2627

/note="L1M4 repeat: matches 5635..5810 of consensus"

2628..2935

/note="AluSq repeat: matches 1..308 of consensus"

2936..3001

/note="L1M4 repeat: matches 5569..5635 of consensus"

3002..3305

/note="AluJb repeat: matches 1..300 of consensus"

3306..3764

/note="L1M4 repeat: matches 5120..5569 of consensus"

3765..4077

/note="AluJo repeat: matches 1..292 of consensus"

4078..4223

/note="L1M4 repeat: matches 4973..5120 of consensus"

4224..4361

/note="AluJo/FPAM repeat: matches 158..294 of consensus"

/complement(join(4666..4827,5364..5428,7817..7945))

/gene="dJ828H9.2"

/product="dJ828H9.2 (putative novel transcript)"

/note="match: ESTs: Em:A1218423"

/evidence=not experimental

/complement(4666)

/gene="dJ828H9.2"

polyA\_site

repeat\_region

5065..5364

/note="AluSx repeat: matches 1..298 of consensus"

5519..5794

/note="L2 repeat: matches 2405..2680 of consensus"

5934..6244

/note="match: GSS: Em:AQ415067"

6021..6321

/note="match: GSS: Em:AQ531024"

6258..7406

/note="L1P4S repeat: matches 4995..6144 of consensus"

8230..8343

/note="WIR repeat: matches 45..166 of consensus"

/complement(join(9193..9766,11907..12175))

/gene="dJ828H9.1"

/complement(join(9193..9766,11907..12175))

/gene="dJ828H9.1"

/product="dJ828H9.1 (putative novel transcript)"

/note="match: ESTs: Em:AA431369 Em:AA382348 Em:AA397834 Em:AA758864 Em:AA906363 Em:AA757279 Em:AW341461 Em:AA432366"

/evidence=not experimental

/complement(9193)

/gene="dJ828H9.1"

/complement(9426)

/gene="dJ828H9.1"

10546..10649

/note="L2 repeat: matches 541..654 of consensus"

10708..10784

/note="ORSL repeat: matches 206..275 of consensus"

10785..11076

/note="AluY repeat: matches 2..295 of consensus"

11077..11230

/note="ORSL repeat: matches 275..458 of consensus"

/complement(12059..12555)

/note="match: GSS: Em:AQ276946"

13524..13668

/note="MER5B repeat: matches 17..166 of consensus"

14312..14516

/note="AluJb repeat: matches 111..306 of consensus"

14691..15441

/note="L1MB3 repeat: matches 5264..6016 of consensus"

15414..15494

/note="L1ME3A repeat: matches 6025..6104 of consensus"

15495..15805

/note="AluSq repeat: matches 1..309 of consensus"

15976..16160

/note="WIRIG repeat: matches 39..229 of consensus"

16253..16343

/note="MER5A repeat: matches 58..145 of consensus"

16275..16389

/note="MER5A repeat: matches 1..123 of consensus"

16406..16912

/note="match: GSS: Em:AQ792876"

16434..16597

/note="WIR repeat: matches 91..262 of consensus"

17457..17911

/note="match: GSS: Em:AQ135214"

17487..17994

/note="match: GSS: Em:AQ215088"

17672..18210

/note="MER52A repeat: matches 1207..1755 of consensus"

18311..18921

/note="MER52A repeat: matches 660..1349 of consensus"

18976..19568

/note="MER4-internal repeat: matches 4586..5102 of consensus"

19578..24868

/note="L1P4S repeat: matches 10..5050 of consensus"

24869..25168

/note="AluSq repeat: matches 1..300 of consensus"

25169..26305

/note="L1P4S repeat: matches 5050..6163 of consensus"

26308..26385



```

/note="MER4-internal repeat: matches 3313..3393 of
consensus"
repeat_region
26386..26686
/note="AluJb repeat: matches 1..302 of consensus"
repeat_region
26687..26832
/note="MER4-internal repeat: matches 3393..3543 of
consensus"
repeat_region
26833..27129
/note="AluXx repeat: matches 1..296 of consensus"
repeat_region
27130..27559
/note="MER4-internal repeat: matches 3543..3989 of
consensus"
repeat_region
27560..27865
/note="AluJb repeat: matches 1..301 of consensus"
repeat_region
27866..28012
/note="MER4-internal repeat: matches 3989..4140 of
consensus"
repeat_region
28013..28262
/note="AluJb repeat: matches 3..264 of consensus"
repeat_region
28263..28514
/note="MER4-internal repeat: matches 4140..4401 of
consensus"
repeat_region
28515..28797
/note="AluXx repeat: matches 1..284 of consensus"
repeat_region
28798..29064
/note="MER4-internal repeat: matches 4401..6347 of
consensus"
misc_feature
28993..29239

```

## Alignment Scores:

```

Pred. No.: 353 Length: 167013
Score: 9.00 Matches: 9
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 45.00% Indels: 0
DB: 9 Gaps: 0

```

US-10-799-747-116 (1-20) x HSDJ828H9 (1-167013)

Qy 5 ServalleuSerPheLeuLeuTyrThr 13

Db 84047 TCTGTGCTCAGCTTCTTCTCTGGACA 84021

## RESULT 9

AC122809/c

LOCUS

DEFINITION Mus musculus BAC clone RP23-272E5 from chromosome 7, complete

sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Mus musculus

Bukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.

1 (bases 1 to 205725)

Swearngen-Shahid, S. and Meyer, R.

The sequence of Mus musculus BAC clone RP23-272E5

Unpublished (2001)

2 (bases 1 to 205725)

Wilson, R.

Sequencing of Mus musculus

Unpublished (2001)

3 (bases 1 to 205725)

McPherson, J.D. and Waterston, R.H.

Direct Submission

Submitted (25-MAY-2002) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

4 (bases 1 to 205725)

McPherson, J.D. and Waterston, R.H.

Direct Submission

Submitted (13-FEB-2003) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

## REFERENCE

## AUTHORS

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## AUTHORS

## TITLE

## JOURNAL

## COMMENT

5 (bases 1 to 205725)

Wilson, R.K.

Direct Submission

Submitted (29-JUL-2003) Genome Sequencing Center, 4444 Forest Park

Parkway, St. Louis, MO 63108, USA

6 (bases 1 to 205725)

Wilson, R.

Direct Submission

Submitted (25-NOV-2003) Department of Genetics, Washington

University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

On Jul 29, 2003 this sequence version replaced gi:28372763.

----- Genome Center

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: <http://genome.wustl.edu>

Contact: [submissions@watson.wustl.edu](mailto:submissions@watson.wustl.edu)

----- Summary Statistics

-----

Center project name: M\_BA0272E05

-----

NOTICE: This sequence may not represent the entire insert of this

clone. It may be shorter because we only sequence overlapping

clone sections once, or longer because we provide a small overlap

between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:

all regions were double stranded, sequenced with an alternate

chemistry, or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats; all regions were covered by sequence

from more than one subclone; and the assembly was confirmed by

restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. Wes Warren,

Department of Genetics, Washington University, St. Louis MO. For

additional information about the map position of this sequence, see

<http://genome.wustl.edu>

SOURCE INFORMATION:

The RPECI-23 BAC Library has been constructed by Kazutoyo Osegawa

and Minako Tateno in the laboratory of Pieter de Jong

(<http://www.chori.org>) from female C57BL/6J mouse kidney and/or

brain genomic DNA. The clone and detailed information can be

obtained from Research Genetics, Inc. (<http://www.resgen.com>) or

Pieter de Jong and coworkers at <http://www.chori.org>

NEIGHBORING SEQUENCE INFORMATION:

This sequence is the entire insert of the clone.

Location/Qualifiers

1..205725

/organism="Mus musculus"

/mol\_type="genomic DNA"

/db\_xref="taxon:10090"

/chromosome="7"

/map="7"

/clone="RP23-272E5"

/clone\_lib="RPECI-23"

1..94

/rpt\_family="Alu"

92..106

/rpt\_family="MaLR"

107..295

/rpt\_family="B2"

296..525

/rpt\_family="MaLR"

706..778

/rpt\_family="Alu"

831..960

/rpt\_family="MIR"

1158..1200

/rpt\_family="Alu"

1201..1596

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

repeat\_region

```

repeat_region /rpt_family="MaLR"
1597.1667 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
1817.1930 /rpt_family="L1"
repeat_region /rpt_family="Alu"
2015.2069 /rpt_family="ID"
repeat_region /rpt_family="ID"
2694.2809 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
2866.3093 /rpt_family="B2"
repeat_region /rpt_family="B2"
3155.3231 /rpt_family="ERV1"
repeat_region /rpt_family="Alu"
3302.3431 /rpt_family="Alu"
repeat_region /rpt_family="L1"
4114.4317 /rpt_family="L1"
repeat_region /rpt_family="L1"
4320.4395 /rpt_family="L1"
repeat_region /rpt_family="Alu"
4396.4542 /rpt_family="Alu"
repeat_region /rpt_family="L1"
4543.4609 /rpt_family="L1"
repeat_region /rpt_family="Alu"
4610.4759 /rpt_family="Alu"
repeat_region /rpt_family="L1"
4760.4898 /rpt_family="L1"
repeat_region /rpt_family="L1"
4899.5280 /rpt_family="MaLR"
repeat_region /rpt_family="L1"
5281.6123 /rpt_family="L1"
repeat_region /rpt_family="L1"
6129.6149 /rpt_family="MaLR"
repeat_region /rpt_family="Alu"
6150.6253 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
6340.6478 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
8595.8732 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
8626.8823 /rpt_family="MaLR"
repeat_region /rpt_family="MaLR"
9217.9562 /rpt_family="MaLR"
repeat_region /rpt_family="MaLR"
1555.1569 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
15734.15835 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
16956.17076 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
18004.18146 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
18164.18345 /rpt_family="B2"
repeat_region /rpt_family="B2"
18848.18921 /rpt_family="ID"
repeat_region /rpt_family="ID"
18938.19051 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
19472.19645 /rpt_family="B2"
repeat_region /rpt_family="B2"
19689.19795 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
20808.20887 /rpt_family="L1"
repeat_region /rpt_family="L1"
21305.21329 /rpt_family="B4"
repeat_region /rpt_family="B4"
21330.21417 /rpt_family="Alu"
repeat_region /rpt_family="Alu"
21436.21596 /rpt_family="MaLR"
repeat_region /rpt_family="MaLR"
21557.21757 /rpt_family="MaLR"
repeat_region /rpt_family="MaLR"
21771.21812 /rpt_family="Alu"

```

```

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repeat_region 22448.22581 /rpt_family="L1"
repeat_region 23288.23405 /rpt_family="Alu"
repeat_region 23447.23515 /rpt_family="ID"
repeat_region 23839.24523 /rpt_family="EMER19A"
repeat_region 24541.24659 /rpt_family="L1"
repeat_region 24690.24764 /rpt_family="ID"
repeat_region 25308.25445 /rpt_family="Alu"
repeat_region 26207.26322 /rpt_family="Alu"
repeat_region 27425.27598 /rpt_family="B4"
repeat_region 27620.27673 /rpt_family="ID"
repeat_region 27674.27706 /rpt_family="B4"
repeat_region 27703.27841 /rpt_family="B2"

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## Alignment Scores:

Pred. No.:	426	Length:	205725
Score:	9.00	Matches:	9
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	45.00%	Indels:	0
DB:	10	Gaps:	0

US-10-799-747-116 (1-20) x AC122809 (1-205725)

QY 3 AlaHisserValLeuSerPheLeuLeu 11  
 |||||  
 Db 136760 GCACATTCAGTTTGTCTATTTTGCTA 136734

## RESULT 10

AC110315 231768 bp DNA linear HTG 09-NOV-2002  
 Rattus norvegicus clone CH230-191N21, \*\*\* SEQUENCING IN PROGRESS  
 DEFINITION \*\*\* 2 unordered pieces.  
 AC110315  
 AC110315.5 GI:24818671  
 HTG; HTGS\_PHASE1; HTGS\_DRAFT; HTGS\_ENRICHED.  
 KEYWORDS Rattus norvegicus (Norway rat)  
 SOURCE Rattus norvegicus  
 ORGANISM Rattus norvegicus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
 Rattus.

## REFERENCE

## AUTHORS

1 (bases 1 to 231768)  
 Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
 Anyalebechi, V., Ayodeji, A., Ayodeji, M., Baca, E., Baden, H.,  
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,  
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,  
 Bryant, N., Buhay, C., Burch, P., Burrell, K., K., Calderon, E.,  
 Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Chen, Z., Chu, J.,  
 Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,  
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,  
 Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,  
 Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K.,  
 Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K.,  
 Egan, A., Escotto, M., Eugene, C., Evans, C. A., Falls, T., Fan, G.,  
 Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P.,  
 Fraser, C. M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M.,  
 Gebregorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W.,  
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Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hoques, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhinney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczek, R., Wooden, H., Worley, K., Wright, D., Wright, J., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A.

## TITLE

## JOURNAL

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## REFERENCE

## AUTHORS

## TITLE

## JOURNAL

## COMMENT

Direct Submission

Unpublished

2 (bases 1 to 231768)

Worley, K.C.

Direct Submission

Submitted (11-FEB-2002) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 231768)

Direct Submission

Rat Genome Sequencing Consortium.

Submitted (09-NOV-2002) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Nov 9, 2002 this sequence version replaced gi:23265946.

The sequence in this assembly is a combination of BAC based reads

and whole genome shotgun sequencing reads assembled using Atlas

(<http://www.hgsc.bcm.tmc.edu/projects/rat/>). Each contig described

in the feature table below represents a scaffold in the Atlas

assembly (a 'contig-scaffold'). Within each contig-scaffold,

individual sequence contigs are ordered and oriented, and separated

by sized gaps filled with Ns to the estimated size. The sequence

may extend beyond the ends of the clone and there may be sequence

contigs within a contig-scaffold that consist entirely of whole

genome shotgun sequence reads. Both end sequences and whole genome

shotgun sequence only contigs will be indicated in the feature

table.

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: GOGP

Center clone name: CH230-191N21

----- Summary Statistics

Assembly program: Phrap; version 0.990329

Consensus quality: 218739 bases at least Q40

Consensus quality: 220995 bases at least Q30

Consensus quality: 22405 bases at least Q20

Estimated insert size: 223718; sum-of-contigs estimation

Quality coverage: 6x in Q20 bases; sum-of-contigs estimation

-----

\* NOTE: Estimated insert size may differ from sequence length  
 (see [http://www.hgsc.bcm.tmc.edu/docs/genbank\\_draft\\_data.html](http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html)).  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 2 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

\* 1 230654: contig of 230654 bp in length

\* 230655 230754: gap of unknown length

\* 230755 231768: contig of 1014 bp in length.

## FEATURES

## source

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/mol\_type="genomic DNA"

/db\_xref="taxon:10116"

/clone="CH230-191N21"

## misc\_feature

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/note="clone\_boundary"

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site:EcoRI

end\_sequence:BH282813"

complement(222634..223294)

/note="clone\_boundary"

clone\_end:T7

site:EcoRI

end\_sequence:BH282812"

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/note="wgs\_end\_extension"

clone\_end:T7"

228778..230654

/note="wgs\_end\_extension"

clone\_end:T7"

## ORIGIN

## Alignment Scores:

Pred. No.:

Score: 475

Percent Similarity: 9.00

Best Local Similarity: 100.00%

Query Match: 45.00%

Indels: 2

Gaps: 0

Length: 231768

Matches: 9

Conservative: 0

Mismatches: 0

Indels: 0

Gaps: 0

US-10-799-747-116 (1-20) x AC110315 (1-231768)

QY 6 ValLeuSerPheLeuLeuTriThrPro 14

|||||

Db 172304 GTGTTTCCTCCCTGTTATGACTCCA 172330

|||||

RESULT 11

AC103176/c

LOCUS

AC103176 254839 bp DNA linear HTG 13-MAY-2003

DEFINITION

Rattus norvegicus clone CH230-98N18, WORKING DRAFT SEQUENCE, 4

unordered pieces.

AC103176

AC103176.5 GI:30578547

HTG; HTGS PHASE1; HTGS DRAFT; HTGS\_FULLTOP.

KEYWORDS

Rattus norvegicus (Norway rat)

SOURCE

ORGANISM

Rattus norvegicus

Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;

Rattus.

1 (bases 1 to 254839)

Muzny, D., Marie, Metzker, M., Lee, S., Adams, C., Alder, J.,

Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,

Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,

Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,

Biswalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, J., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregorjis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hognes, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhea, L., Louiseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapa, P., Martin, K., Martin, R., Martinez, E., Mashiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwakoileme, O., Okwuonu, G., Olarnpungagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plapper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puaor, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rivers, C., Rodkey, T., Rojas, A., Rose, R., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Soza, J., Steinle, M., Strong, R., Sutton, A., Svatok, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, P., Williams, G., Willson, R., Wleczkyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausen, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstein, G., and Gibbs, R.A.  
 Direct Submission  
 Unpublished  
 2 (bases 1 to 254839)  
 Worley, K.C.  
 Direct Submission  
 Submitted (24-NOV-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
 3 (bases 1 to 254839)  
 Rat Genome Sequencing Consortium.  
 Direct Submission  
 Submitted (13-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA  
 On May 13, 2003 this sequence version replaced gi:23265304.  
 The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.

Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: http://www.hgsc.bcm.tmc.edu/  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information  
 Center project name: GJKE  
 Center clone name: CH230-98N18  
 ----- Summary Statistics  
 Assembly program: Atlas 3.0;  
 Consensus quality: 241531 bases at least Q40  
 Consensus quality: 243998 bases at least Q30  
 Consensus quality: 246135 bases at least Q20  
 Estimated insert size: 252438; sum-of-contigs estimation  
 Quality coverage: 7x in Q20 bases; sum-of-contigs estimation

\* NOTE: Estimated insert size may differ from sequence length  
 \* (see http://www.hgsc.bcm.tmc.edu/docs/genbank\_draft\_data.html)  
 \* NOTE: This sequence may represent more than one clone.  
 \* NOTE: This is a 'working draft' sequence. It currently  
 \* consists of 4 contigs. The true order of the pieces  
 \* is not known and their order in this sequence record is  
 \* arbitrary. Gaps between the contigs are represented as  
 \* runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 \* as soon as it is available and the accession number will  
 \* be preserved.

\* 1 250539: contig of 250539 bp in length  
 \* 250540 250839: gap of unknown length  
 \* 250640 251762: contig of 1123 bp in length  
 \* 251763 251862: gap of unknown length  
 \* 251863 252935: contig of 1073 bp in length  
 \* 252936 253035: gap of unknown length  
 \* 253036 254839: contig of 1804 bp in length.

## FEATURES

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 /organism="Rattus norvegicus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:10116"  
 /clone="CH230-98N18"

## misc\_feature

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 clone end:T7"  
 3845..5494

## misc\_feature

/note="wgs end\_extension  
 clone end:T7"  
 complement(6002..29990)

## misc\_feature

/note="clone boundary  
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 site:EcoRI

end\_sequence:BH311760"

complement(247726..248554)

/note="clone boundary  
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## ORIGIN

## Alignment Scores:

Pred. No.: 518 Length: 254839  
 Score: 9.00 Matches: 9  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 45.00% Indels: 0  
 DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC103176 (1-254839)

Qy 5 ServValLeuSerPheLeuLeuTrpThr 13

Db 107271 TCTGTACTTAGTTTCTCTGTGGACT 107245

RESULT 12

AC098074/c

----- Genome Center

LOCUS AC098074 267352 bp DNA linear HTG 10-MAY-2003  
 DEFINITION Rattus norvegicus clone CH230-119124, WORKING DRAFT SEQUENCE, 4  
 AC098074  
 ACCESSION AC098074.7 GI:30522668  
 VERSION HTG; HTGS\_PHASE1; HTGS DRAFT; HTGS\_FULLTOP.  
 KEYWORDS Rattus norvegicus (Norway rat)  
 SOURCE Rattus norvegicus  
 ORGANISM Rattus norvegicus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.  
 1 (bases 1 to 267352)  
 REFERENCE Muzny, D. Marie., Metzker, M. Lee., Abramson, S., Adams, C., Alder, J.,  
 Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,  
 Anyalibechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H.,  
 Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F.,  
 Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M.,  
 Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E.,  
 Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A.,  
 Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J.,  
 Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L.,  
 Davila, M. L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D.,  
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 Lorensuewa, L., Lounseged, H., Lorado, R. J., Lu, X., Ma, J.,  
 Maheshwari, M., Mahindaratne, M., Mamoud, M., Malloy, K., Mangum, A.,  
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 Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L.,  
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 Pasternak, S., Paul, H., Perez, A., Perez, L., Pfankuch, C.,  
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 Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F.,  
 Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K.,  
 Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V.,  
 Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von  
 Niederhausern, A., Weiss, R., Smith, D. R., Holt, R. A., Smith, H. O.,  
 Weinstock, G., and Gibbs, R. A.  
 Direct Submission  
 Unpublished  
 2 (bases 1 to 267352)  
 REFERENCE Worley, K. C.  
 TITLE Direct Submission  
 JOURNAL  
 Submitted (23-OCT-2001) Human Genome Sequencing Center, Department  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 3 (bases 1 to 267352)  
 REFERENCE Rat Genome Sequencing Consortium.  
 TITLE Direct Submission  
 JOURNAL Submitted (10-MAY-2003) Human Genome Sequencing Center, Department

COMMENT  
 of Molecular and Human Genetics, Baylor College of Medicine, One  
 Baylor Plaza, Houston, TX 77030, USA  
 On May 10, 2003 this sequence version replaced gi:24635703.  
 The sequence in this assembly is a combination of BAC based reads  
 and whole genome shotgun sequencing reads assembled using Atlas  
 (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described  
 in the feature table below represents a scaffold in the Atlas  
 assembly (a 'contig-scaffold'). Within each contig-scaffold,  
 individual sequence contigs are ordered and oriented, and separated  
 by sized gaps filled with Ns to the estimated size. The sequence  
 may extend beyond the ends of the clone and there may be sequence  
 contigs within a contig-scaffold that consist entirely of whole  
 genome shotgun sequence reads. Both end sequences and whole genome  
 shotgun sequence only contigs will be indicated in the feature  
 table.  
 ----- Genome Center  
 Center: Baylor College of Medicine  
 Center code: BCM  
 Web site: http://www.hgsc.bcm.tmc.edu/  
 Contact: hgsc-help@bcm.tmc.edu  
 ----- Project Information  
 Center project name: GSGO  
 Center clone name: CH230-119124  
 ----- Summary Statistics  
 Assembly program: Atlas 3.0;  
 Consensus quality: 254595 bases at least Q40  
 Consensus quality: 257284 bases at least Q30  
 Consensus quality: 259179 bases at least Q20  
 Estimated insert size: 267168; sum-of-contigs estimation  
 Quality coverage: 7x in Q20 bases; sum-of-contigs estimation  
 -----  
 \* NOTE: Estimated insert size may differ from sequence length  
 (see http://www.hgsc.bcm.tmc.edu/docs/genbank\_draft\_data.html)  
 \* NOTE: This sequence may represent more than one clone.  
 \* NOTE: This is a 'working draft' sequence. It currently  
 consists of 4 contigs. The true order of the pieces  
 is not known and their order in this sequence record is  
 arbitrary. Gaps between the contigs are represented as  
 runs of N, but the exact sizes of the gaps are unknown.  
 \* This record will be updated with the finished sequence  
 as soon as it is available and the accession number will  
 be preserved.  
 \* 1 242526: contig of 242526 bp in length  
 \* 242527 242626: gap of unknown length  
 \* 242627 264340: contig of 21714 bp in length  
 \* 264341 264440: gap of unknown length  
 \* 264441 265515: contig of 1075 bp in length  
 \* 265516 267352: gap of unknown length  
 \* 267352 267352: contig of 1737 bp in length.  
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 source  
 1. 267352  
 /organism="Rattus norvegicus"  
 /mol\_type="genomic DNA"  
 /db\_xref="taxon:10116"  
 /clone="CH230-119124"  
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 /note="clone boundary  
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 site:EcoRI  
 end sequence:BH316141"  
 18571. 19415  
 /note="clone boundary  
 clone end:Sp6  
 site:EcoRI  
 end\_sequence:BH316143"  
 misc\_feature  
 misc\_feature  
 Alignment Scores:  
 Pred. No.: 541 Length: 267352  
 Score: 9.00 Matches: 9  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 45.00% Indels: 0

ORIGIN  
 Alignment Scores:  
 Pred. No.: 541 Length: 267352  
 Score: 9.00 Matches: 9  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 45.00% Indels: 0

DB:	2	Gaps:	0
US-10-799-747-116	(1-20) x AC098074	(1-267352)	
QY	5 SerValLeuSerPheLeuLeuTrpThr	13	
DB	247232 TCTGACTAGTTTCTCTGTGGACT	247206	
RESULT 13			
AC098226/c			
LOCUS	AC098226	306660 bp	DNA linear HTG 13-MAY-2003
DEFINITION	Rattus norvegicus clone CH230-56K13, *** SEQUENCING IN PROGRESS		
ACCESSION	AC098226		
VERSION	HTG; HTGS PHASE1; HTGS DRAFT; HTGS ENRICHED.		
KEYWORDS	Rattus norvegicus (Norway rat)		
SOURCE	Rattus norvegicus		
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.		
REFERENCE	1 (bases 1 to 306660)		
AUTHORS	Muzny, D. Marie, Metzker, M. Lee, Abramson, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswal, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Egan, A., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregorgis, E., Geet, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamill, C., Hamilton, C., Hernandez, J., Harvey, Y., Haylak, P., Hawes, A., Henderson, N., Hernandez, J., Hollins, B., Hines, S., Hladun, S.L., Hodgson, A., Hogue, M., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpach, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowalski, C., Kraft, C.I., Lebowitz, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensu, H., Loulseghe, H., Lozano, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokeme, O., Okunolu, G., Olarnpusagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkuch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savary, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajls, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steinle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villalana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G., and Gibbs, R.A.		
	Direct Submission		
TITLE	Unpublished		
JOURNAL			
REFERENCE	2 (bases 1 to 306660)		
AUTHORS	Worley, K.C.		
JOURNAL	Direct Submission		
	Submitted (23-OCT-2001) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA		
REFERENCE	3 (bases 1 to 306660)		
AUTHORS	Rat Genome Sequencing Consortium.		
JOURNAL	Direct Submission		
	Submitted (13-MAY-2003) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA		
COMMENT	On May 13, 2003 this sequence version replaced gi:23268205. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas ( <a href="http://www.hgsc.bcm.tmc.edu/projects/rat/">http://www.hgsc.bcm.tmc.edu/projects/rat/</a> ). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table.		
	----- Genome Center		
	Center: Baylor College of Medicine		
	Center code: BCM		
	Web site: <a href="http://www.hgsc.bcm.tmc.edu/">http://www.hgsc.bcm.tmc.edu/</a>		
	Contact: hgsc-help@bcm.tmc.edu		
	----- Project Information		
	Center project name: GIVQ		
	Center clone name: CH230-56K13		
	----- Summary Statistics		
	Assembly program: Atlas 3.0;		
	Consensus quality: 236654 bases at least Q40		
	Consensus quality: 241334 bases at least Q30		
	Consensus quality: 24452 bases at least Q20		
	Estimated insert size: 240001; sum-of-contigs estimation		
	Quality coverage: 6x in Q20 bases; sum-of-contigs estimation		
	-----		
	* NOTE: Estimated insert size may differ from sequence length		
	(see <a href="http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html">http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html</a> ).		
	* NOTE: This is a 'working draft' sequence. It currently		
	* consists of 11 contigs. The true order of the pieces		
	* is not known and their order in this sequence record is		
	* arbitrary. Gaps between the contigs are represented as		
	* runs of N, but the exact sizes of the gaps are unknown.		
	* This record will be updated with the finished sequence		
	* as soon as it is available and the accession number will		
	* be preserved.		
	-----		
	1 4465: contig of 4465 bp in length		
	4466 4565: gap of unknown length		
	4566 280032: contig of 275467 bp in length		
	280033 280132: gap of unknown length		
	280133 291663: contig of 11531 bp in length		
	291664 291763: gap of unknown length		
	291764 297095: contig of 5332 bp in length		
	297096 297195: gap of unknown length		
	297196 298558: contig of 1363 bp in length		
	298559 298659: gap of unknown length		
	298659 299805: contig of 1147 bp in length		
	299806 301125: gap of unknown length		
	301126 301225: gap of unknown length		
	301226 302302: contig of 1077 bp in length		
	302303 302402: gap of unknown length		
	302403 303604: contig of 1202 bp in length		
	303605 303704: gap of unknown length		
	303705 305546: contig of 1842 bp in length		
	305547 305646: gap of unknown length		
	305647 306660: contig of 1014 bp in length.		
	Location/Qualifiers		
FEATURES			

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1. .306660
/organism="Rattus norvegicus"
/mol_type="genomic DNA"
/db_xref="taxon:10116"
/clone="CH230-56K13"
misc_feature
1. .1130
/note="wgs end extension"
clone_end:Sp6"
misc_feature
4566. .5998
/note="wgs end extension"
clone_end:Sp6"
misc_feature
29817. .30488
/note="clone boundary"
clone_end:Sp6"
site:EcoRI
end sequence:BH272628"
misc_feature
31694. .33434
/note="wgs contig"
misc_feature
159947. .161767
/note="wgs contig"
complement(265403. .266254)
/note="clone boundary"
clone_end:T7
site:EcoRI
end sequence:BH272626"
misc_feature
272770. .274704
/note="wgs end extension"
clone_end:T7"
misc_feature
278461. .280032
/note="wgs end extension"
clone_end:T7"

ORIGIN
Alignment Scores:
Pred. No.: 613 Length: 306660
Score: 9.00 Matches: 9
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 45.00% Indels: 0
DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AC098226 (1-306660)

Qy 6 ValLeuSerPheLeuLeuTrpThrPro 14
Db 215746 GTGCTTTCTCTCTGTTATGGACTCCA 215720

RESULT 14
G25673
LOCUS human STS EST113435, sequence tagged site. 339 bp DNA linear STS 02-JUN-1996
DEFINITION human STS EST113435, sequence tagged site.
ACCESSION G25673
VERSION G25673.1 GI:1347905
KEYWORDS STS; STS sequence; primer; sequence tagged site.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Hudson, T.
REFERENCE 1 (bases 1 to 339)
AUTHORS Whitehead Institute/MIT Center for Genome Research; Physically
TITLES Mapped STSs
JOURNAL Unpublished (1995)
COMMENT
Contact: Thomas Hudson
Whitehead Institute/MIT Center for Genome Research
9 Cambridge Center, Cambridge MA 02142 USA
Tel: 617 252 1900
Fax: 617 252 1902
Email: thudson@genome.wi.mit.edu

Primer A: AATTACAGTTTGGTTCTCAGATT
Primer B: CCATGCTAAGTCCAGTGGGG

```

```

Primer B: CCATGCTAAGTCCAGTGGGG
STS size: 150
PCR Profile:
  Presoak:
    Denaturation:
      Annealing: 56 degrees C
    Polymerization:
      PCR Cycles: 35
    Thermal Cycler:
      Template: 10 ng
      Primer: each 5 pM
      dNTPs: each 4 nM
      Taq Polymerase: 0.025 units/ul
      Total Vol: 20 ul
  Buffer:
    MgCl2: 1.5 mM
    KCl: 50 mM
    Tris-HCl: 10 mM
    pH: 9.3
Derived from dbEST (genbank accession T59368).
FEATURES
  source
    1..339
    /organism="Homo sapiens"
    /mol_type="genomic DNA"
    /db_xref="taxon:9606"
    /map="527.4 cR from top of Chr2 linkage group"
  STS
    primer_bind
      1..150
    primer_bind
      1..24
      complement(131..150)
  ORIGIN
Alignment Scores:
Pred. No.: 13.1 Length: 339
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 11 Gaps: 0

US-10-799-747-116 (1-20) x G25673 (1-339)

Qy 3 AlaHisSerValLeuSerPheLeu 10
Db 108 GCTCATTCTGTACTTCTCTCTC 131

RESULT 15
G27758
LOCUS human STS SHGC-33168, sequence tagged site. 356 bp DNA linear STS 29-JUN-1996
DEFINITION human STS SHGC-33168, sequence tagged site.
ACCESSION G27758
VERSION G27758.1 GI:11396477
KEYWORDS STS; STS sequence; primer; sequence tagged site.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Myers, R.M.
REFERENCE 1 (bases 1 to 356)
AUTHORS Stanford Human Genome Center (SHGC)
JOURNAL Department of University School of Medicine
COMMENT Unpublished (1996)
Contact: Richard M. Myers
Stanford Human Genome Center (SHGC)
Stanford University School of Medicine
Department of Genetics, M-344, Stanford, CA 94305, USA
Tel: 4157259687
Fax: 4157259689
Email: myers@shgc.stanford.edu

Primer A: AATTACAGTTTGGTTCTCAGATT
Primer B: CCATGCTAAGTCCAGTGGGG

```

STS size: 150

PCR Profile:

Initial incubation: 94 degrees C for 90 seconds

Denaturation: 94 degrees C for 15 seconds

Annealing: 62 degrees C for 23 seconds

Polymerization: 72 degrees C for 30 seconds

PCR Cycles: 30

Thermal Cycler: Perkin Elmer 9600

Protocol:

Template: 25 ng

Primer: each 1 uM

dNTPs: each 200 uM

Taq Polymerase: 0.05 units/ul

Total Vol: 10 ul

Buffer:

MgCl2: 2.5 mM

KCl: 50 mM

Tris-HCl: 20 mM

pH: 8.3

Prepared with primer pairs provided by Sandoz, derived from T59368

-- Washington University/Merck EST sequence.

# FEATURES

source

1..356

/organism="Homo sapiens"

/mol\_type="genomic DNA"

/db\_xref="taxon:9606"

/map="2"

1..150

1..24

complement(131..150)

# ORIGIN

Alignment Scores:

Pred. No.:	13.7	Length:	356
Score:	8.00	Matches:	8
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	40.00%	Indels:	0
DB:	11	Gaps:	0

US-10-799-747-116 (1-20) x G27758 (1-356)

QY 3 AlaHisSerValLeuSerPheLeu 10

Db 108 GCTCATTCTGTACTTTCTTCCTC 131

Search completed: July 21, 2004, 04:43:02

Job time : 1826 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model  
Run on: July 21, 2004, 02:53:21 ; Search time 649 Seconds  
(without alignments)  
130.915 Million cell updates/sec

Title: US-10-799-747-116  
Perfect score: 20  
Sequence: 1 MAHSVLSFLWTPVAKSX 20

Scoring table: OLIGO  
Xgapop 60.0 , Xgapext 60.0  
Ygapop 60.0 , Ygapext 60.0  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 3373863 seqs, 2124099041 residues  
Word size: 1

Total number of hits satisfying chosen parameters: 6742599

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Command line parameters:  
-MODEL=frame+ p2n.model -DEV=xlp  
-Q=/cgn2\_1/USPRO.spool\_p/US10799747/runat\_19072004\_161441\_21967/app\_query.fasta\_1.199  
-DB=N\_Geneseq\_29Jan04 -QMT=fastap -SUFFIX=olig.rng -MINMATCH=0.1 -LOOPCL=0  
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi  
-LIST=45 -DOCALIGN=200 -THR SCORE=quality -THR MIN=1 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=pt0 -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10799747 @CGN 1.1 708 @runat\_19072004\_161441\_21967 -NCPU=6 -ICPU=3  
-NO.MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DEV\_TIMEOUT=120 -WARN\_TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=60  
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database : N\_Geneseq\_29Jan04.\*  
1: Geneseqn1980s.\*  
2: Geneseqn1990s.\*  
3: Geneseqn2000s.\*  
4: Geneseqn2001as.\*  
5: Geneseqn2001bs.\*  
6: Geneseqn2002s.\*  
7: Geneseqn2003as.\*  
8: Geneseqn2003bs.\*  
9: Geneseqn2003cs.\*  
10: Geneseqn2004s.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	19	95.0	1434	2	AAX37452
2	19	95.0	1434	7	ADA39771 Human sec
3	19	95.0	1434	7	ACC50424 Human sec
4	8	40.0	356	6	ABL77790 Human ova
5	8	40.0	1503	7	ADA70841
6	7	35.0	49	7	ACD97100 Rice gene
7	7	35.0	65	6	ABZ28939 Candida g
8	7	35.0	90	4	AAS23646 Tetracycl

9	7	35.0	109	2	AAH86006
10	7	35.0	155	7	ACD97494
c 11	7	35.0	263	2	AAT20076
12	7	35.0	264	9	ADC93542
c 13	7	35.0	293	6	ABN76057
c 14	7	35.0	294	9	ADC92876
c 15	7	35.0	315	2	AAV88416
c 16	7	35.0	352	6	ABQ85808
c 17	7	35.0	363	5	ABV25515
c 18	7	35.0	377	5	AAE66743
c 19	7	35.0	379	8	ACH50509
c 20	7	35.0	394	4	ACH87798
c 21	7	35.0	409	5	ABV38500
c 22	7	35.0	415	3	AAH30357
c 23	7	35.0	426	5	ABV08598
c 24	7	35.0	432	6	ABK76313
c 25	7	35.0	433	7	ABX61964
c 26	7	35.0	436	4	ACH32772
c 27	7	35.0	440	8	ACH32772
c 28	7	35.0	445	8	ACH42384
c 29	7	35.0	455	8	ACH15424
c 30	7	35.0	465	4	AAI11211
31	7	35.0	465	4	AAI11211
32	7	35.0	465	4	AAI32473
33	7	35.0	465	4	AAK67520
34	7	35.0	465	4	ABA42442
35	7	35.0	465	4	ABA22654
36	7	35.0	465	4	AAK26581
37	7	35.0	465	4	AAK01123
38	7	35.0	465	4	ABS26173
39	7	35.0	465	5	AAI01128
40	7	35.0	466	6	ABN94789
41	7	35.0	467	8	ACH26710
42	7	35.0	468	4	AAK32950
43	7	35.0	468	4	AAK07199
44	7	35.0	468	4	ABS32679
45	7	35.0	468	4	ABS32679

## ALIGNMENTS

RESULT 1  
AAX37452

ID AAX37452 standard; cDNA; 1434 BP.

XX AAX37452;

DT 06-JUL-1999 (first entry)

XX Human secreted protein cDNA fragment containing gene 2.

Human; secreted protein; treatment; prevention; protein therapy; AIDS; gene therapy; diagnosis; cancer; tumor; neurodegenerative disorder; developmental abnormality; fetal deficiency; blood disorder; leukemia; immune system disease; autoimmune disease; hepatic disease; lymphoma; renal disease; inflammation; allergy; Alzheimer's disease; schizophrenia; cognitive disorder; prostate disease; skeletal; cardiac; muscle disorder; pulmonary disorder; transplant rejection; osteoclast; osteoporosis; arthritis; malignancy; digestive; endocrine; infection; ss.

XX Homo sapiens.

XX WO9918208-A1.

PN 15-APR-1999.

XX 01-OCT-1998; 98MO-US020775.

XX 02-OCT-1997; 97US-0060833P.

PR 02-OCT-1997; 97US-0060836P.

PR 02-OCT-1997; 97US-0060837P.

PR 02-OCT-1997; 97US-0060838P.

PR	02-OCT-1997;	97US-0060843P.	
PR	02-OCT-1997;	97US-0060862P.	
PR	02-OCT-1997;	97US-0060866P.	
PR	02-OCT-1997;	97US-0060874P.	
PR	02-OCT-1997;	97US-0060880P.	
PR	02-OCT-1997;	97US-0060884P.	
XX		(HUMA-) HUMAN GENOME SCI INC.	
PA			
XX	Duan DR, Florence KA, Rosen CA, Ruben SM, Greene JM, Young P;		
PI	Ferrie AM, Yu G, Janat F, Ni J, Carter KC, Endress GA, Peng P;		
PI	Lafleur DW, Shi Y;		
XX			
DR	WPI; 1999-264022/22.		
DR	P-PSDB; AAY07853.		
XX			
PI	New isolated human genes and the secreted polypeptides they encode.		
XX			
PS	Claim 1a; Page 228; 368pp; English.		
XX			
CC	This invention describes novel isolated human genes and the secreted		
CC	proteins they encode. The products of the invention are useful for		
CC	preventing, treating or ameliorating medical conditions, e.g. by protein		
CC	or gene therapy. Also pathological conditions can be diagnosed by		
CC	determining the amount of the new polypeptides in a sample or by		
CC	determining the presence of mutations in the new polynucleotides.		
CC	Specific uses are described for each of the 101 polynucleotides, based on		
CC	which tissues they are most highly expressed in, and include developing		
CC	products for the diagnosis or treatment of cancer, tumours,		
CC	neurodegenerative disorders, developmental abnormalities and fetal		
CC	deficiencies, blood disorders, leukemias, diseases of the immune system,		
CC	autoimmune diseases, hepatic and renal disease, lymphomas, inflammation,		
CC	allergies, Alzheimer's and cognitive disorders, schizophrenia, prostate		
CC	disease, skeletal or cardiac muscle disorders, pulmonary disorders,		
CC	transplant rejection, disorders involving osteoclasts such as		
CC	osteoporosis, arthritis or malignancies, digestive/endocrine disorders,		
CC	infections and AIDS. The human secreted proteins of the invention are		
CC	represented in AAY07852-Y07993 and the encoding nucleic acids are		
CC	represented in AAX37451-X37552		
XX			
SQ	Sequence 1434 BP; 480 A; 204 C; 250 G; 495 T; 0 U; 5 Other;		
Alignment Scores:			
Pred. No.:	5,49e-10	Length:	1434
Score:	19.00	Matches:	19
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	95.00%	Indels:	0
DB:	2	Gaps:	0
US-10-799-747-116 (1-20) x AAX37452 (1-1434)			
QY	1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19		
Db	507 ATGGCAGCCCAATTCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTCTGAATCA 563		
RESULT 2			
ADA39771			
ID	ADA39771 standard; cDNA; 1434 BP.		
XX			
AC	ADA39771;		
XX			
DT	20-NOV-2003 (first entry)		
XX			
DE	Human secreted protein encoding cDNA.		
XX			
KW	Human; secreted protein; cancer; hyperproliferative disorder;		
KW	rheumatoid arthritis; autoimmune disorder; haematopoietic disorder;		
KW	anaemia; allergic reaction; asthma; cardiovascular disorder;		
KW	wound healing; cytostatic; immunosuppressive; neurotropic; neuroprotective;		
KW	antiviral; antiallergic; hepatotropic; antidiabetic; antiinflammatory;		
KW	vulnery; cardiant; gene therapy; ss.		
XX			

OS	Homo sapiens.		
XX			
FN	WO2002102993-A2.		
XX			
PD	27-DEC-2002.		
XX			
PF	19-MAR-2002; 2002WO-US008123.		
XX			
PR	21-MAR-2001; 2001US-0277340P.		
PR	19-JUL-2001; 2001US-0306171P.		
PR	13-NOV-2001; 2001US-0331287P.		
XX			
PA	(HUMA-) HUMAN GENOME SCI INC.		
XX			
PI	Rosen CA, Ruben SM;		
XX			
DR	WPI; 2003-175238/17.		
XX			
PT	New human secreted proteins and nucleic acid molecules, useful for		
PT	preparing a diagnostic or pharmaceutical composition for diagnosing,		
PT	preventing or treating cancer or other hyperproliferative disorder,		
XX	asthma, allergies or AIDS.		
PS	Claim 9; SEQ ID NO 153; 3205pp; English.		
XX			
CC	The invention relates to novel genes ADA39629-ADA40565 and proteins		
CC	ADA40566-ADA41501 for human secreted proteins, useful for preventing,		
CC	treating or ameliorating medical conditions e.g. by protein or gene		
CC	therapy. The polypeptides, nucleic acid molecules, antibodies or their		
CC	fragments, and agonists or antagonists that bind to the polypeptide are		
CC	useful for preparing a diagnostic or pharmaceutical composition for		
CC	diagnosing or treating cancer or other hyperproliferative disorder. The		
CC	polypeptides and nucleic acid molecules are also useful for detecting,		
CC	preventing, diagnosing, prognosticating, treating or ameliorating cancer		
CC	or other hyperproliferative disorders including neoplasms, autoimmune		
CC	disorders (e.g. diabetes, rheumatoid arthritis, systemic lupus		
CC	erythematosus, multiple sclerosis, autoimmune thyroiditis or haemolytic		
CC	anaemia), haematopoietic or haematological disorders (e.g. anaemia,		
CC	thrombocytopenia), allergic reactions including asthma or eczema,		
CC	inflammatory disorders (e.g. ischaemia-reperfusion injury, inflammatory		
CC	bowel disease or Crohn's disease), neurodegenerative disorders (e.g.		
CC	Alzheimer's disease or Parkinson's disease), cardiovascular disorders		
CC	(e.g. atherosclerosis, myocarditis), infectious diseases (bacterial,		
CC	fungal or viral infections including HIV/AIDS), or wound healing and		
CC	disorders of epithelial cell proliferation. The nucleic acids are also		
CC	useful for chromosome identification, radiation hybrid mapping or long-		
CC	range restriction mapping, as molecular weight markers, or as		
CC	hybridization or diagnostic probes. The polypeptides and antibodies are		
CC	useful for providing immunological probes for differential identification		
CC	of the tissues immunohistochemistry assays. Note: The sequence data for		
CC	this patent did not form part of the printed specification, but was		
CC	obtained in electronic format directly from WIPO at		
CC	ftp.wipo.int/pub/published_pct_sequences.		
XX			
SQ	Sequence 1434 BP; 480 A; 203 C; 250 G; 496 T; 0 U; 5 Other;		
Alignment Scores:			
Pred. No.:	5,49e-10	Length:	1434
Score:	19.00	Matches:	19
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	95.00%	Indels:	0
DB:	7	Gaps:	0
US-10-799-747-116 (1-20) x ADA39771 (1-1434)			
QY	1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyrAlaLeuLysSer 19		
Db	507 ATGGCAGCCCAATTCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTCTGAATCA 563		
RESULT 3			
ACC50424			
ID	ACC50424 standard; cDNA; 1434 BP.		



Wed Jul 21 09:11:28 2004

RESULT 5  
ADA70841  
ID ADA70841 standard; DNA; 1503 BP.  
XX  
AC ADA70841;  
XX  
DT 20-NOV-2003 (first entry)  
XX  
DE Rice gene, SEQ ID 4164.  
XX  
DE Plant; bacterial infection; fungal infection; viral infection; rice;  
KW gene; ds.  
XX  
OS Oryza sativa.  
XX  
PN WO2003000898-A1.  
XX  
PD 03-JAN-2003.  
XX  
PF 22-JUN-2001; 2001WO-IB001105.  
XX  
PR 22-JUN-2001; 2001WO-IB001105.  
XX  
PA (SYGN ) SYNGENTA PARTICIPATIONS AG.  
XX  
PI Chang H, Chen W, Cooper B, Glazebrook J, Goff SA, Hou Y;  
PI Katagiri F, Quan S, Tao Y, Whitham S, Xie Z, Zhu T, Zou G;  
XX  
DR WPI; 2003-175290/17.  
XX  
PT Identifying at least one gene involved in plant resistance or response to  
PT pathogenic infection for conferring resistance or tolerance to a plant to  
PT bacterial, fungal or viral infection by determining or detecting plant  
PT gene expression.  
XX  
PS Claim 6; SEQ ID NO 4164; 899pp; English.  
XX  
CC The present invention relates to a method (M1) for identifying genes  
CC involved in plant resistance or response to pathogenic infection. M1  
CC comprises identifying a gene whose expression is significantly altered in  
CC the incompatible interaction of plant gene expression relative to  
CC the expression of the gene in an uninfected plant, in a mutant plant that  
CC does not express a gene associated with response to pathogenic infection,  
CC or in a corresponding incompatible or compatible interaction. (M1) is  
CC useful for conferring resistance to resistance or tolerance to a plant to  
CC bacterial, fungal or viral infection. The present sequence was used to  
CC illustrate the invention.  
XX  
SQ Sequence 1503 BP; 395 A; 329 C; 383 G; 396 T; 0 U; 0 Other;

Alignment Scores:  
Pred. No.: 72.6 Length: 1503  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 7 Gaps: 0

US-10-799-747-116 (1-20) x ADA70841 (1-1503)  
QY 3 AlaHisSerValIleuSerPheLeu 10  
DB 266 GCACACAGTGTATCTTTCCT 289  
RESULT 6  
ACD97100/c  
ID ACD97100 standard; cDNA; 49 BP.  
XX  
AC ACD97100;  
XX  
DT 23-SEP-2003 (first entry)  
XX

Human colon cancer cell expressed cDNA #5512.  
Open reading frame detection; genome sequencing; colon cancer;  
breast cancer; population genome analysis; genetic shift; cancer;  
antibiotic resistance; antibiotic non-tolerance; congenital disease;  
agriculture; food crop genome; resistance gene; retrovirus;  
influenza virus; eukaryotic pathogen detection; trypanosome; Plasmodium;  
gene; ss.  
Homo sapiens.  
US2002155438-A1.  
24-OCT-2002.  
27-SEP-1999; 99US-00406117.  
20-NOV-1998; 98US-00196716.  
(SIMP/) SIMPSON A J G.  
(NETO/) NETO E D.  
(BREN/) BRENTANI R R.  
Simpson AJG, Neto ED, Brentani RR;  
WPI; 2003-182626/18.  
Determining open reading frames of genome of an organism e.g. a human  
suffering from cancer involves use of single oligonucleotide primer at  
low stringency for preparing single-stranded cDNA from mRNA of  
individual.  
Example 9; Page 787; 959pp; English.  
The invention describes a method of determining open reading frames in  
the genome of organism, comprising contacting mRNA from cell of organism  
with a single oligonucleotide primer (I) at low stringency, preparing  
single-stranded cDNA by reverse transcribing mRNA with (I), amplifying  
cDNA, sequencing the product, and repeating the contacting, preparing  
and amplifying steps with different primers and sequencing resulting  
nucleic acids. The method is useful for: determining that a known  
nucleotide sequence from a genome of an organism corresponds to a  
nucleotide sequence of an open reading frame; for preparing a contig,  
nucleic acid molecule from a genome of an organism; and for sequencing  
all or part of a genome of an organism. mRNA is obtained from mammalian  
or human cell which is associated with a pathological condition e.g. a  
colon cancer or breast cancer cell. The method is useful for analyses of  
populations of subjects and can be used to carry out genetic analyses of  
large or small populations. further, it can be used to study living  
systems to determine if, e.g. there have been genetic shifts which render  
an individual or population more or less likely to be afflicted with  
diseases such as cancer, to determine antibiotic resistance or non-  
tolerance, and so forth. The method can also be used in the study of  
congenital diseases, and the risk of affliction to a foetus, as well as  
the study of whether the conditions are likely to be passed to offspring  
through ova or sperm. The analyses for pathological conditions can be  
carried out in all animals, plants, birds, fish, etc. Using this method,  
in the area of agriculture, for example the genomes of food crops can be  
studied to determine if resistance genes are present, defects in plant  
genomes can also be studied in this way. Similarly, the method permits  
determination of the pathogens which integrate into the genome, such as  
retroviruses and other integrating viruses such as influenza virus, have  
undergone shifts or mutations, which may require different approaches to  
therapy. This method is also applied to eukaryotic pathogens, such as  
trypanosomes, different types of Plasmodium, etc. The method essentially  
eliminates sequencing of non-coding portions. This sequence represents a  
polynucleotide isolated from human colon cancer cell cDNA library  
Sequence 49 BP; 20 A; 10 C; 10 G; 6 T; 0 U; 3 Other;

Alignment Scores:  
Pred. No.: 31.2 Length: 49  
Score: 7.00 Matches: 7

Percent Similarity: 100.00%  
 Best Local Similarity: 100.00%  
 Query Match: 35.00%  
 DB: 7  
 Gaps: 0  
 Indels: 0  
 Mismatches: 0  
 Conservative: 0

US-10-799-747-116 (1-20) x ACD97100 (1-49)

OY 6 ValLeuSerPheLeuLeuTrp 12  
 DB 32 GTTTAAGCTTCTCTCTTTG 12

## RESULT 7

ABZ28939  
 ID ABZ28939 standard; DNA; 65 BP.  
 XX  
 AC ABZ28939;  
 XX  
 DT 30-JAN-2003 (first entry)  
 XX  
 DE Candida gene related tetracycline promoter PCR primer SEQ ID NO 3022.  
 XX  
 KW Fungus; yeast; tetracycline; promoter; GRACE strain; biosynthesis;  
 KW signal transduction; DNA replication; cell division; growth;  
 KW proliferation; Candida albicans; fungicide; antifungal; PCR; primer; ss.  
 XX  
 OS Candida albicans.  
 XX  
 FN WO200253728-A2.  
 XX  
 PD 11-JUL-2002.  
 XX  
 PF 26-DEC-2001; 2001WO-US049486.  
 XX  
 PR 29-DEC-2000; 2000US-0259128P.  
 PR 20-FEB-2001; 2001US-00792024.  
 PR 22-AUG-2001; 2001US-0314050P.  
 XX  
 PA (ELIT-) ELITRA PHARM INC.  
 XX  
 PI Roemer T, Jiang B, Boone C, Bussey H, Ohlsen KL;  
 XX  
 DR WPI; 2002-566694/60.  
 XX  
 PS Constructing strains for identifying gene products as effective targets  
 CC for therapeutic intervention, by inactivating in the strain one allele of  
 CC a gene and placing other allele of the gene under conditional expression.  
 CC  
 CC Claim 76; SEQ ID NO 3022; 167pp + Sequence Listing; English.

The invention relates to constructing (M1) a strain of diploid fungal cells in which both alleles of a gene are modified, comprising modifying one allele by insertion or replacement by a cassette having an expressible selectable marker and modifying other allele by recombination, of a promoter replacement fragment with a heterologous promoter, so that expression of the second allele is regulated by the promoter. (M1) is useful for constructing a strain of diploid fungal cells in which both alleles of a gene are modified. The diploid fungal cells having both alleles modified are useful for identifying a gene that is essential to the survival or growth of a fungus, a gene that contributes to the virulence and/or pathogenicity of a fungus, a gene that contributes to the resistance of a diploid fungus to an antifungal agent, an antifungal agent that inhibits the growth of a diploid fungus and for identifying a therapeutic agent for treatment of a diploid fungus disease. (M1) is useful for identifying a compound which modulates the activity of a gene product, preferably enzymatic activity, carbon compound catabolism, biosynthetic, transporter, transcriptional, translational, signal transduction, DNA replication and cell division activity. The method is useful for identifying a compound having the ability to inhibit growth or proliferation of C. albicans cells and for treating infection by C. albicans. The present sequence is that of a PCR primer used in the method of the invention. Note: The sequence data for this patent is not represented in the printed specification but is based on sequence information supplied to Derwent by the European Patent Office

XX  
 SQ Sequence 65 BP; 17 A; 12 C; 10 G; 26 T; 0 U; 0 Other;

## Alignment Scores:

Pred. No.: 40.5 Length: 65  
 Score: 7.00 Matches: 7  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 35.00% Indels: 0  
 DB: 6 Gaps: 0

US-10-799-747-116 (1-20) x ABZ28939 (1-65)

OY 5 SerValLeuSerPheLeuLeu 11  
 DB 13 TCTGTATTATCATCTCTTTATG 33

## RESULT 8

AAS23646  
 ID AAS23646 standard; DNA; 90 BP.  
 XX  
 AC AAS23646;  
 XX  
 DT 04-DEC-2001 (first entry)  
 XX  
 DE Tetracycline promoter downstream PCR primer (Tet-Down) #21.  
 XX  
 KW Gene identification; essential gene; GRACE; pathogenic fungus;  
 KW gene replacement and conditional expression; fungal infection;  
 KW PCR primer; Tet-Down; tetracycline promoter; ss.  
 XX  
 OS Escherichia coli.  
 OS Candida albicans.  
 XX  
 PN WO200160975-A2.  
 XX  
 PD 23-AUG-2001.  
 XX  
 PF 20-FEB-2001; 2001WO-US005551.  
 XX  
 PR 18-FEB-2000; 2000US-0183534P.  
 XX  
 PA (ELIT-) ELITRA PHARM INC.  
 XX  
 PI Roemer T, Jiang B, Boone C, Bussey H;  
 XX  
 DR WPI; 2001-489080/53.  
 XX  
 PS Identifying genes essential to fungal metabolisms and identifying  
 CC potential therapeutic agents that target these genes.  
 CC  
 CC Disclosure; Page 292; 324pp; English.

The present invention relates to novel methods for constructing fungal strains useful for identification and validation of gene products as targets for therapeutic agents, for creating a collection of identified essential genes, and screening assays for the discovery of new drugs. The invention provides the GRACE (gene replacement and conditional expression) method for the construction of mutant organisms referred to as GRACE strains of the organism. The invention can be applied to any organism, particularly a pathogenic fungus e.g. Candida albicans, Aspergillus fumigatus and Cryptococcus neoformans. The methods are useful to identify agents that may be used in the treatment of fungal infections. AAS23626-AAS23686 represent tetracycline promoter downstream PCR primers (Tet-Down) used in the methods of the present invention

SQ Sequence 90 BP; 23 A; 15 C; 17 G; 35 T; 0 U; 0 Other;

## Alignment Scores:

Pred. No.: 54.7 Length: 90  
 Score: 7.00 Matches: 7  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0

Query Match:	35.00%	Indels:	0
DB:	4	Gaps:	0
US-10-799-747-116 (1-20) x AAS23646 (1-90)			
Qy	5 SerValleuSerPheLeuLeu 11		
Db	13 TCTGATTATCATCTTATTG 33		
RESULT 9			
AAH86006			
ID	AAH86006 standard; DNA; 109 BP.		
AC			
XX	AAH86006;		
XX			
DT	27-FEB-2002 (first entry)		
XX			
DE	Human single nucleotide polymorphism containing DNA sequence #863.		
XX			
KW	Biallelic marker; polymorphism; human; disease; diagnosis; treatment;		
KW	phenotypic trait; gene therapy; forensic; paternity; mapping; cancer;		
KW	transgenic; single nucleotide polymorphism; SNP; ss.		
XX			
OS	Homo sapiens.		
XX			
FH	Key	Location/Qualifiers	
FT	variation	replace(47,A)	
FT		/*tag= a	
FT		/standard_name= "single nucleotide polymorphism"	
XX			
PN	WO9953095-A2.		
XX			
PD	21-OCT-1999.		
XX			
XX	30-MAR-1999; 99WO-US006893.		
XX			
PR	09-APR-1998; 98US-00057871.		
XX			
XX	(WHED ) WHITEHEAD INST BIOMEDICAL RES.		
PA			
XX			
PI	Lander ES, Wang D, Hudson T;		
XX			
DR	WPI; 1999-620443/53.		
XX			
PT	Polymorphic human genomic sequences and related allele-specific probes		
PT	and primers, useful for genetic analysis, e.g. diagnosis and monitoring		
PT	of disease.		
XX			
PS	Claim 1; Page 123; 330pp; English.		
XX			
CC	This invention describes novel human nucleic acid segments (I) containing		
CC	polymorphic sites. The polynucleotides of (I) are used for, e.g.		
CC	correlating disease polymorphisms (or disease susceptibility) or other		
CC	phenotypic traits (e.g. baldness, obesity, fertility, strength, response		
CC	to drugs etc.); diagnosing and monitoring e.g. cancer, inflammation,		
CC	heart or central nervous system diseases; detecting susceptibility to		
CC	microbial infection; treating or preventing such diseases; forensic		
CC	analysis; gene therapy; paternity testing; mapping genomic loci		
CC	associated with phenotypic traits (and subsequent cloning of the genes		
CC	responsible); and the production of transgenic organisms. Antibodies		
CC	raised against (I) are useful as diagnostic and therapeutic tools and in		
CC	drug screening. AAH85144 - AAH87644 represent the human DNA sequences		
XX	containing biallelic polymorphic sites described in the invention		
XX			
SQ	Sequence 109 BP; 30 A; 28 C; 18 G; 33 T; 0 U; 0 Other;		
Alignment Scores:			
Pred. No.:	65.4	Length:	109
Score:	7.00	Matches:	7
Percent Similarity:	100.00%	Conservative:	0
Best Local Similarity:	100.00%	Mismatches:	0
Query Match:	35.00%	Indels:	0
DB:	2	Gaps:	0

US-10-799-747-116 (1-20) x AAH86006 (1-109)	
Qy	5 SerValleuSerPheLeuLeu 11
Db	15 AGTGTACTGTCTTTTCTACTG 35
RESULT 10	
ACD97494	
ID	ACD97494 standard; cDNA; 155 BP.
XX	
AC	ACD97494;
XX	
DT	23-SEP-2003 (first entry)
XX	
DE	Human colon cancer cell expressed cDNA #5906.
XX	
KW	Open reading frame detection; genome sequencing; colon cancer;
KW	breast cancer; population genome analysis; genetic shift; cancer;
KW	antibiotic resistance; antibiotic non-tolerance; congenital disease;
KW	agriculture; food crop genome; resistance gene; retrovirus;
KW	influenza virus; eukaryotic pathogen detection; trypanosome; Plasmodium;
KW	gene; ss.
XX	
OS	Homo sapiens.
XX	
FN	US2002155438-A1.
XX	
PD	24-OCT-2002.
XX	
PF	27-SEP-1999; 99US-00406117.
XX	
PR	20-NOV-1998; 98US-00196716.
XX	
PA	(SIMP/) SIMPSON A J G.
PA	(NETO/) NETO E D.
PA	(BREN/) BRENTANI R R.
XX	
PI	Simpson AUG, Neto ED, Brentani RR;
XX	
DR	WPI; 2003-182626/18.
XX	
PT	Determining open reading frames of genome of an organism e.g. a human
PT	suffering from cancer involves use of single oligonucleotide primer at
PT	low stringency for preparing single-stranded cDNA from mRNA of
PT	individual.
XX	
PS	Example 9; Page 842; 959pp; English.
XX	
CC	The invention describes a method of determining open reading frames in
CC	the genome of organism, comprising contacting mRNA from cell of organism
CC	with a single oligonucleotide primer (I) at low stringency, preparing
CC	single-stranded cDNA by reverse transcribing mRNA with (I), amplifying
CC	cDNA, sequencing the product, and repeating the contacting, preparing
CC	and amplifying steps with different primers and sequencing resulting
CC	nucleic acids. The method is useful for: determining that a known
CC	nucleotide sequence from a genome of an organism corresponds to a
CC	nucleic acid molecule from a genome of an organism; and for sequencing
CC	all or part of a genome of an organism. mRNA is obtained from mammalian
CC	or human cell which is associated with a pathological condition e.g. a
CC	colon cancer or breast cancer cell. The method is useful for analyses of
CC	populations of subjects and can be used to carry out genetic analyses of
CC	large or small populations. further, it can be used to study living
CC	systems to determine if, e.g. there have been genetic shifts which render
CC	an individual or population more or less likely to be afflicted with
CC	diseases such as cancer, to determine antibiotic resistance or non-
CC	tolerance, and so forth. The method can also be used in the study of
CC	congenital diseases, and the risk of affliction to a foetus, as well as
CC	the study of whether the conditions are likely to be passed to offspring
CC	through ova or sperm. The analyses for pathological conditions can be
CC	carried out in all animals, plants, birds, fish, etc. Using this method,
CC	in the area of agriculture, for example the genomes of food crops can be

CC studied to determine if resistance genes are present, defects in plant  
 CC genomes can also be studied in this way. Similarly, the method permits  
 CC determination of the pathogens which integrate into the genome, such as  
 CC retroviruses and other integrating viruses such as influenza virus, have  
 CC undergone shifts or mutations, which may require different approaches to  
 CC therapy. This method is also applied to eukaryotic pathogens, such as  
 CC trypanosomes, different types of plasmidium, etc. The method essentially  
 CC eliminates sequencing of non-coding portions. This sequence represents a  
 CC polynucleotide isolated from human colon cancer cell cDNA library

XX SQ Sequence 155 BP; 41 A; 30 C; 33 G; 49 T; 0 U; 2 Other;

Alignment Scores: Pred. No.: 90.6 Length: 155  
 Score: 7.00 Matches: 7  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 35.00% Indels: 0  
 DB: 7 Gaps: 0

US-10-799-747-116 (1-20) x ACD97494 (1-155)

Qy 5 SerValLeuSerPheLeuLeu 11  
 |||||  
 Db 12 AGTGTGCTTCTTCTCTTA 32

RESULT 11

AAT20076/c  
 ID AAT20076 standard; cDNA to mRNA; 263 BP.

XX AC AAT20076;

DT 17-JUL-1996 (first entry)

XX Human gene signature HUMGS01218.

XX Gene signature; messenger RNA; mRNA; relative abundance; frequency;  
 XX human; cloning; mapping; non-biased library; diagnosis; detection;  
 XX cell typing; abnormal cell function; ss.

XX Homo sapiens.

XX WO9514772-A1.

XX 01-JUN-1995.

XX 11-NOV-1994; 94WO-JP001916.

XX 12-NOV-1993; 93JP-00355504.

XX (MATS/) MATSUBARA K.  
 XX (OKUB/) OKUBO K.

XX Matsubara K, Okubo K;

XX MPI; 1995-206931/27.

XX Single-stranded DNA for identifying gene signatures - isolated from 3'-  
 XX directed human cDNA library that reflects relative abundance of corresp.  
 XX mRNA in specific human tissues.

XX Claim 1; Page 554; 2245pp; Japanese.

XX A single-stranded DNA (or its complementary strand or the corresp. double  
 XX -stranded DNA) which comprises one of the 7837 "GS" sequences given in  
 XX AAT19001-T26837 and which is able to hybridise to part of human genomic  
 XX DNA, cDNA or mRNA is claimed. The GS (Gene Signature) sequences were  
 XX obtained from 3'-directed cDNA libraries prepared from various human  
 XX tissues; synthesis of cDNA was initiated from the 3'-end of mRNA by using  
 XX poly(T) as the sole primer. Since the 3'- untranslated sequence is unique  
 XX to a particular mRNA species, almost all the 3'-oriented cDNAs hybridise  
 XX with specific mRNAs. Each library is constructed so as to reflect  
 XX accurately the relative abundance of different mRNAs in the particular

CC tissue from which it was derived. The appearance frequency of a given GS  
 CC in a cDNA library can be determined (esp. using primers and probes  
 CC derived from the GS sequences) as a means of diagnosing abnormal cell  
 CC function or for recognising different cell types

XX SQ Sequence 263 BP; 92 A; 34 C; 43 G; 92 T; 0 U; 2 Other;

Alignment Scores: Pred. No.: 148 Length: 263  
 Score: 7.00 Matches: 7  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 35.00% Indels: 0  
 DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AAT20076 (1-263)

Qy 5 SerValLeuSerPheLeuLeu 11  
 |||||  
 Db 82 AGTGTGCTTCTTCTCTC 62

RESULT 12

ADC93542  
 ID ADC93542 standard; DNA; 264 BP.

XX AC ADC93542;

XX 01-JAN-2004 (first entry)

XX E. faecium DNA sequence SEQ ID 3169.

XX ds; gene; urinary tract infection; bacteraemia; endocarditis; wound;  
 XX abdominal-pelvic infection.

XX Enterococcus faecium.

XX US6583275-B1.

XX 24-JUN-2003.

XX 30-JUN-1998; 98US-00107532.

XX 02-JUL-1997; 97US-0051571P.

XX 14-MAY-1998; 98US-0085598P.

XX (GENO-) GENOME THERAPEUTICS CORP.

XX Doucette-Stamm LA, Rush D;

XX WPI; 2003-799836/75.

XX P-PSDB; ADC97196.

XX New isolated nucleic acid derived from Enterococcus faecium encoding an  
 XX Enterococcus faecium polypeptide useful for detection, prevention and  
 XX treatment of a pathological condition resulting from a bacterial  
 XX infection.

XX Example 1; SEQ ID NO 3169; 243pp; English.

XX The invention relates to an isolated nucleic acid derived from  
 XX Enterococcus faecium encoding an Enterococcus faecium polypeptide having  
 XX one of 10 fully defined sequences given in the (or comprising 40  
 XX sequential nucleotides chosen from any of the nucleic acids, its  
 XX complement or sequences hybridising to it). Also included are a  
 XX recombinant vector comprising the nucleic acid operably linked to  
 XX transcription regulatory element, a cell comprising the vector and a  
 XX single-stranded probe comprising the nucleic acid. The nucleic acids are  
 XX chosen from 3654 disclosed sequences encoding 3654 disclosed proteins.  
 XX The nucleic acids is useful for diagnosing pathological conditions  
 XX resulting from E. faecium bacterial infection (e.g. urinary tract  
 XX infection, bacteraemia, endocarditis, wounds and abdominal-pelvic  
 XX infection) and for screening drugs such as agonists and antagonists. The  
 XX nucleic acid is useful for recombinant production of Candida albicans -

CC derived peptides or antisense polypeptides. Pharmaceutical compositions  
CC and vaccines containing the nucleic acid are useful for preventing or  
CC treating Enterococcus faecium infections. The present sequence represents  
CC one of the disclosed E. faecium nucleic acids.

SQ Sequence 264 BP; 67 A; 59 C; 48 G; 88 T; 0 U; 2 Other;

Alignment Scores:  
Pred. No.: 148 Length: 264  
Score: 7.00 Matches: 7  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 35.00% Indels: 0  
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x ADC93542 (1-264)

Qy 5 SerValLeuSerPheLeuLeu 11  
|||||  
Db 104 TCGGTCTCGTCTTCTCTTTA 124

RESULT 13

ABN76057/c  
ID ABN76057 standard; cDNA; 293 BP.

AC ABN76057;

DT 08-JUL-2002 (first entry)

DE Human ORF1004 cDNA, SEQ ID NO:2007.

XX Human; ORF; open reading frame; ORFX; drug screening; diagnosis;  
KW disease monitoring; cytokine; cell proliferation; cell differentiation;  
KW immune modulation; haematopoiesis regulation; tissue growth;  
KW angiogenesis; activin; inhibin; chemotactic; chemokinetic; haemostatic;  
KW thrombolytic; tumour inhibition; bodily characteristic; fertility;  
KW behaviour; cancer; proliferative disorder; neurological disorder;  
KW cardiovascular disease; immune system disorder; organ transplantation;  
KW tissue growth disorder; tissue regeneration disorder; diabetes mellitus;  
KW hypothyroidism; cholesterol ester storage disease; infection; vulnery;  
KW vasotropic; antipsoriatic; antidiabetic; cytostatic; nootropic;  
KW neuroprotective; antithrombotic; anticoagulant; thrombolytic;  
KW cardiatic; hypotensive; antithyroid; antiinflammatory; immunomodulator;  
KW dermatological; analgesic; virucide; antibacterial; fungicide; gene; ss.

XX Homo sapiens.

XX WO200190366-A2.

PN 29-NOV-2001.

PD 24-MAY-2001; 2001WO-US017076.

PF 24-MAY-2000; 2000US-0206690P.

PR (CURA-) CUPAGEN CORP.

PA Leach MD, Shinkets RA;

PI WPI; 2002-106200/14.

XX P-PSDB; ABP32031.

XX Novel human polypeptides and polynucleotides useful for diagnosing,  
PT preventing and treating cardiovascular disease, neurodegenerative,  
PT hyperproliferative disorders and disorders related to organ  
PT transplantation.

XX Claim 1; Page 766; 2508pp; English.

XX Sequences ABP31028-ABP35561 represent 4534 novel human proteins  
CC designated ORF (open reading frame) 1-4534, and sequences ABN75054-  
CC ABN79587 represent cDNAs encoding them. The invention also encompasses  
CC polypeptides at least 80% identical to the ORF1-ORF4534 (collectively

CC referred to as ORFX) proteins, polynucleotides at least 85% identical to  
CC the ORFX nucleic acid sequences, vectors and host cells comprising ORFX  
CC polynucleotides, the recombinant production of ORFX proteins, antibodies  
CC specific for ORFX proteins, methods of detecting ORFX polynucleotides and  
CC polypeptides, methods of screening for modulators of ORFX expression or  
CC activity, and methods of screening individuals for a predisposition to an  
CC ORFX-associated disorder. The ORFX proteins of the invention have a wide  
CC range of biological activities, such as cytokine, cell proliferation,  
CC cell differentiation, immune modulation, haematopoiesis regulation,  
CC tissue growth, angiogenesis, activin or inhibin activity, chemotactic/  
CC chemokinetic activity, haemostatic activity, thrombolytic activity,  
CC receptor/ligand, antiinflammatory activity, tumour inhibition activity,  
CC and antiinfective activity, and may also be involved in the determination  
CC of bodily characteristics, fertility and behaviour. ORFX proteins,  
CC nucleic acids and antibodies may be used in the treatment of cancers,  
CC other proliferative disorders such as psoriasis and benign tumours,  
CC neurological disorders such as epilepsy and Alzheimer's disease,  
CC cardiovascular diseases, immune system disorders, disorders related to  
CC organ transplantation, disorders of tissue growth and regeneration,  
CC diseases such as diabetes mellitus, hypothyroidism, and cholesterol ester  
CC storage disease, and infectious diseases caused by viral, bacterial,  
CC fungal and other pathogens. ORFX nucleic acids may also be used as a  
CC source of primers and probes, in the detection of ORFX genomic sequences  
CC or transcripts, in the identification and cloning of homologous  
CC sequences, in genetic diagnosis, and in forensic biology. The ORFX  
CC nucleic acids may additionally be used to produce transgenic animals  
CC which may be useful for studying the function and/or activity of ORFX  
CC protein, and in drug screening. The ORFX proteins may also be used as  
CC immunogens to generate specific antibodies, which are useful in the  
CC diagnosis, treatment and monitoring of ORFX-associated diseases

XX Sequence 293 BP; 74 A; 85 C; 66 G; 66 T; 0 U; 2 Other;

Alignment Scores:

Pred. No.: 163 Length: 293  
Score: 7.00 Matches: 7  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 35.00% Indels: 0  
DB: 6 Gaps: 0

US-10-799-747-116 (1-20) x ABN76057 (1-293)

Qy 5 SerValLeuSerPheLeuLeu 11  
|||||  
Db 158 AGTGTCTCAGCTTCTCTT 138

RESULT 14

ADC92876/c

ID ADC92876 standard; DNA; 294 BP.

XX ADC92876;

XX 01-JAN-2004 (first entry)

DE E. faecium DNA sequence SEQ ID 2503.

XX ds; gene; urinary tract infection; bacteraemia; endocarditis; wound;

XX abdominal-pelvic infection.

XX Enterococcus faecium.

XX US6583275-B1.

PN 24-JUN-2003.

XX 30-JUN-1998; 98US-00107532.

XX 02-JUL-1997; 97US-0051571P.

PR 14-MAY-1998; 98US-0085598P.

XX (GENO-) GENOME THERAPEUTICS CORP.

XX



```

PI Doucette-Stamm LA, Bush D;
XX
XX WPI: 2003-799836/75.
DR P-PSDB; ADC96530.
XX
PT New isolated nucleic acid derived from Enterococcus faecium encoding an
PT Enterococcus faecium polypeptide useful for detection, prevention and
PT treatment of a pathological condition resulting from a bacterial
PT infection.
XX
XX Example 1; SEQ ID NO 2503; 243pp; English.
PS
XX The invention relates to an isolated nucleic acid derived from
CC Enterococcus faecium encoding an Enterococcus faecium polypeptide having
CC one of 10 fully defined sequences given in the (or comprising 40
CC sequential nucleotides chosen from any of the nucleic acids, its
CC complement or sequences hybridising to it). Also included are a
CC recombinant vector comprising the nucleic acid operably linked to
CC transcription regulatory element, a cell comprising the vector and a
CC single-stranded probe comprising the nucleic acid. The nucleic acids are
CC chosen from 3654 disclosed sequences encoding 3654 disclosed proteins.
CC The nucleic acids are useful for diagnosing pathological conditions
CC resulting from E. faecium bacterial infection (e.g. urinary tract
CC infection, bacteraemia, endocarditis, wounds and abdominal-pelvic
CC infection) and for screening drugs such as agonists and antagonists. The
CC nucleic acid is useful for recombinant production of Candida albicans -
CC derived peptides or antisense polypeptides. Pharmaceutical compositions
CC and vaccines containing the nucleic acid are useful for preventing or
CC treating Enterococcus faecium infections. The present sequence represents
CC one if the disclosed E. faecium nucleic acids.
XX
XX Sequence 294 BP; 106 A; 57 C; 61 G; 68 T; 0 U; 2 Other;

Alignment Scores:
Pred. No.: 164 Length: 294
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x ADC92876 (1-294)

Oy 5 SerValLeuSerPheLeuLeu 11
Db 64 TCGGTTCTGCTTCTCCCTTTA 44

RESULT 15
AAV88416/c
ID AAV88416 standard; cDNA; 315 BP.
XX
XX AAV88416;
AC
XX 12-FEB-1999 (first entry)
DT
DE EST clone EX581.
DE
XX
XX Expressed sequence tag; secreted protein; haematopoiesis regulator;
KW tissue growth; activin; inhibin; tumour invasion suppressor; EST; human;
KW chemotaxis; chemokinesis; haemostasis; gene therapy; thrombolysis;
KW receptor; ligand; anti-inflammatory; tumour inhibitor; ds.
XX
XX Homo sapiens.
OS
XX
XX WO9845437-A2.
PN
XX 15-OCT-1998.
PD
XX
XX 10-APR-1998; 98WO-US006956.
PF
XX 10-APR-1997; 97US-00837312.
PR
XX (GEMY ) GENETICS INST INC.
PA

```

Jacobs K, Mccoy JM, Lavallie ER, Racie LA, Merberg D, Treacy M;  
Spaulding V, Agostino MJ;  
WPI: 1999-070078/06.

New polynucleotides encoding human secreted proteins - derived from e.g.  
human blood, kidney, foetal lung, placenta, testes, brain, ovary,  
pituitary, retina and colon cDNA libraries.

Claim 1; Page 389; 64pp; English.

The present sequence represents an expressed sequence tag (EST), and is a  
polynucleotide of the invention. The polynucleotides of the invention are  
all secreted EST sequences isolated from a variety of human tissue  
sources. The EST sequences and proteins encoded by them are predicted to  
have useful biological activities which would make them suitable for  
treating, preventing or ameliorating medical conditions in humans and  
animals, although no supporting data is given. Suggested activities  
include nutritional activity, immune stimulating or suppressing activity,  
haematopoiesis regulating activity, tissue growth activity, haemostatic  
activity/inhibin activity, chemotactic/chemokinetic activity, haemostatic  
and thrombolytic activity, receptor/ligand activity, anti-inflammatory  
activity, cadherin/tumour invasion suppressor activity, tumour inhibition  
activity. The EST sequences are also stated to be useful for gene therapy

Sequence 315 BP; 74 A; 79 C; 65 G; 97 T; 0 U; 0 Other;

Alignment Scores:  
Pred. No.: 175 Length: 315  
Score: 7.00 Matches: 7  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 35.00% Indels: 0  
DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x AAV88416 (1-315)

Oy 2 AlaAlaHisSerValLeuSer 8  
Db 215 GCAGCACACAGTGTCCTTTCT 195

Search completed: July 21, 2004, 04:12:23  
Job time : 660 secs

***This Page Blank (uspto)***

GenCore version 5.1.1.6  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 03:55:26 ; Search time 72 Seconds  
(without alignments)  
154.153 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 20  
Sequence: 1 MAHSVLSFLWTYPALKSX 20

Scoring table: OLIGO  
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Ygapop 60.0 , Ygapext 60.0  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 682709 seqs, 277475446 residues

Word size: 1

Total number of hits satisfying chosen parameters: 1359535

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Command line parameters:

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-DB=Issued Patents NA -QMT=fastap -SUFFIX=olig.rni -MINMATCH=0.1 -LOOPCL=0  
-LOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=oligo -TRANS=human40.cdi  
-LIST=45 -DOCALIGN=200 -THR SCORE=quality -THR MIN=1 -ALIGN=15 -MODE=LOCAL  
-OUTFMT=ptc -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10799747@cgn\_1\_105@runat\_19072004\_161443\_22019 -NCPU=6 -ICPU=3  
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6  
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database : Issued Patents NA.\*

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- 2: /cgn2\_6/ptodata/2/ina/5B.COMB.seq.\*
- 3: /cgn2\_6/ptodata/2/ina/6A.COMB.seq.\*
- 4: /cgn2\_6/ptodata/2/ina/6B.COMB.seq.\*
- 5: /cgn2\_6/ptodata/2/ina/PCTUS.COMB.seq.\*
- 6: /cgn2\_6/ptodata/2/ina/backfiles.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	7	35.0	264	4	US-09-107-532A-3169
2	7	35.0	294	4	US-09-107-532A-2503
3	7	35.0	894	4	US-09-107-532A-1065
4	7	35.0	933	4	US-09-308-003-46
5	7	35.0	1983	1	US-08-221-817-21
6	7	35.0	1983	1	US-08-454-439-21
7	7	35.0	1983	5	PCT-US94-10487-21
8	7	35.0	2056	2	US-08-836-442-1
9	7	35.0	2560	4	US-09-786-256C-29
10	7	35.0	2569	4	US-08-956-171E-74
11	7	35.0	3024	2	US-08-836-943-1
12	7	35.0	3465	4	US-09-134-000C-2717

C 13	7	35.0	3892	3	US-08-569-214-1	Sequence 1, Appli
C 14	7	35.0	3892	3	US-08-937-236-1	Sequence 1, Appli
C 15	7	35.0	4218	4	US-09-081-385-8	Sequence 8, Appli
C 16	7	35.0	34185	4	US-09-545-481-3	Sequence 3, Appli
C 17	7	35.0	99916	4	US-09-816-095-3	Sequence 3, Appli
C 18	7	35.0	536165	4	US-09-214-808-1	Sequence 1, Appli
C 19	7	35.0	786431	4	US-09-751-389-3	Sequence 3, Appli
C 20	7	35.0	1664976	4	US-08-916-421B-1	Sequence 1, Appli
C 21	7	35.0	1664976	4	US-08-916-421B-1	Sequence 1, Appli
C 22	6	30.0	18	4	US-09-308-003-40	Sequence 40, Appli
C 23	6	30.0	18	4	US-09-422-978-9787	Sequence 9787, Ap
C 24	6	30.0	20	1	US-08-271-942A-86	Sequence 86, Appl
C 25	6	30.0	20	3	US-08-779-916A-86	Sequence 86, Appl
C 26	6	30.0	20	5	PCT-US95-08604-86	Sequence 86, Appl
C 27	6	30.0	29	3	US-08-544-381B-152	Sequence 152, App
C 28	6	30.0	32	3	US-08-575-967A-6	Sequence 6, Appli
C 29	6	30.0	35	3	US-09-282-736-16	Sequence 16, Appl
C 30	6	30.0	40	4	US-10-092-246-28	Sequence 28, Appl
C 31	6	30.0	40	4	US-10-150-051-7	Sequence 7, Appli
C 32	6	30.0	40	4	US-10-150-051-8	Sequence 8, Appli
C 33	6	30.0	45	4	US-08-983-564A-21	Sequence 21, Appl
C 34	6	30.0	47	4	US-09-422-978-2157	Sequence 2157, Ap
C 35	6	30.0	65	2	US-03-121-887-5	Sequence 5, Appli
C 36	6	30.0	65	3	US-09-241-353-5	Sequence 5, Appli
C 37	6	30.0	65	3	US-09-245-984-5	Sequence 5, Appli
C 38	6	30.0	65	3	US-09-241-979-5	Sequence 5, Appli
C 39	6	30.0	107	4	US-09-313-294A-1806	Sequence 1806, Ap
C 40	6	30.0	108	1	US-08-441-591-25	Sequence 25, Appl
C 41	6	30.0	108	1	US-08-303-362A-25	Sequence 25, Appl
C 42	6	30.0	108	5	PCT-US95-05600-42	Sequence 42, Appl
C 43	6	30.0	117	3	US-09-284-832-33	Sequence 33, Appl
C 44	6	30.0	120	3	US-09-284-832-32	Sequence 32, Appl
C 45	6	30.0	138	3	US-09-432-020B-1	Sequence 1, Appli

ALIGNMENTS

RESULT 1

US-09-107-532A-3169  
; Sequence 3169, Application US/09107532A  
; Patent No. 6583275  
; GENERAL INFORMATION:

APPLICANT: Lynn A Doucette-Stamm and David Bush  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS

NUMBER OF SEQUENCES: 7310

CORRESPONDENCE ADDRESS:

ADDRESSEE: GENOME THERAPEUTICS CORPORATION

STREET: 100 Beaver Street

CITY: Waltham

STATE: Massachusetts

COUNTRY: USA

ZIP: 02354

COMPUTER READABLE FORM:

MEDIUM TYPE: CD-ROM ISO9660

COMPUTER: PC

OPERATING SYSTEM: <Unknown>

SOFTWARE: ASCII

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/09/107,532A

FILING DATE: 30-Jun-1998

PRIOR APPLICATION DATA:

APPLICATION NUMBER: 60/085,598

FILING DATE: 14 May 1998

APPLICATION NUMBER: 60/051571

FILING DATE: July 2, 1997

ATTORNEY/AGENT INFORMATION:

NAME: Ariniello, Pamela Deneke

REGISTRATION NUMBER: 40,489

REFERENCE/DOCKET NUMBER: GTC-012

TELECOMMUNICATION INFORMATION:

TELEPHONE: (781)893-5007

TELEFAX: (781)893-8277

INFORMATION FOR SEQ ID NO: 3169:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 264 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Enterococcus faecium  
FEATURE:  
NAME/KEY: misc feature  
LOCATION: (B) LOCATION 1...264  
SEQUENCE DESCRIPTION: SEQ ID NO: 3169:  
US-09-107-532A-3169  
Alignment Scores:  
Pred. No.: 15 Length: 264  
Score: 7.00 Matches: 7  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 35.00% Indels: 0  
DB: 4 Gaps: 0  
US-10-799-747-116 (1-20) x US-09-107-532A-3169 (1-264)  
QY 5 SerValLeuSerPheLeuLeu 11  
DB 104 TCGGTTCTGCTTCTTCCTTTA 124  
RESULT 2  
US-09-107-532A-2503/c  
Sequence 2503, Application US/09107532A  
Patent No. 6583275  
GENERAL INFORMATION:  
APPLICANT: Lynn A Doucette-Stamm and David Bush  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS  
NUMBER OF SEQUENCES: 7310  
CORRESPONDENCE ADDRESS:  
ADDRESS: GENOME THERAPEUTICS CORPORATION  
STREET: 100 Beaver Street  
CITY: Waltham  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 02354  
COMPUTER READABLE FORM:  
MEDIUM TYPE: CD/ROM ISO9660  
COMPUTER: PC  
OPERATING SYSTEM: <Unknown>  
SOFTWARE: ASCII  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/107,532A  
FILING DATE: 30-Jun-1998  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 60/085,598  
FILING DATE: 14 May 1998  
APPLICATION NUMBER: 60/051571  
FILING DATE: July 2, 1997  
ATTORNEY/AGENT INFORMATION:  
NAME: Ariniello, Pamela Deneke  
REGISTRATION NUMBER: 40,489  
REFERENCE/DOCKET NUMBER: GTC-012  
TELEPHONE: (781)893-5007  
TELEFAX: (781)893-8277  
INFORMATION FOR SEQ ID NO: 2503:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 294 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular

MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Enterococcus faecium  
FEATURE:  
NAME/KEY: misc feature  
LOCATION: (B) LOCATION 1...294  
SEQUENCE DESCRIPTION: SEQ ID NO: 2503:  
US-09-107-532A-2503  
Alignment Scores:  
Pred. No.: 16 Length: 294  
Score: 7.00 Matches: 7  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 35.00% Indels: 0  
DB: 4 Gaps: 0  
US-10-799-747-116 (1-20) x US-09-107-532A-2503 (1-294)  
QY 5 SerValLeuSerPheLeuLeu 11  
DB 64 TCGGTTCTGCTTCTTCCTTTA 44  
RESULT 3  
US-09-107-532A-1065/c  
Sequence 1065, Application US/09107532A  
Patent No. 6583275  
GENERAL INFORMATION:  
APPLICANT: Lynn A Doucette-Stamm and David Bush  
TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO  
ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS  
NUMBER OF SEQUENCES: 7310  
CORRESPONDENCE ADDRESS:  
ADDRESS: GENOME THERAPEUTICS CORPORATION  
STREET: 100 Beaver Street  
CITY: Waltham  
STATE: Massachusetts  
COUNTRY: USA  
ZIP: 02354  
COMPUTER READABLE FORM:  
MEDIUM TYPE: CD/ROM ISO9660  
COMPUTER: PC  
OPERATING SYSTEM: <Unknown>  
SOFTWARE: ASCII  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/107,532A  
FILING DATE: 30-Jun-1998  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 60/085,598  
FILING DATE: 14 May 1998  
APPLICATION NUMBER: 60/051571  
FILING DATE: July 2, 1997  
ATTORNEY/AGENT INFORMATION:  
NAME: Ariniello, Pamela Deneke  
REGISTRATION NUMBER: 40,489  
REFERENCE/DOCKET NUMBER: GTC-012  
TELEPHONE: (781)893-5007  
TELEFAX: (781)893-8277  
INFORMATION FOR SEQ ID NO: 1065:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 894 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ANTI-SENSE: NO  
ORIGINAL SOURCE:  
ORGANISM: Enterococcus faecium  
FEATURE:

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; NAME/KEY: misc feature
; LOCATION: (B) LOCATION 1...894
; SEQUENCE DESCRIPTION: SEQ ID NO: 1065:
US-09-107-532A-1065

Alignment Scores:
Pred. No.: 48 Length: 894
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x US-09-107-532A-1065 (1-894)

Qy 5 SerValLeuSerPheLeu 11
Db 887 TCTGTTCTTAGCTTTTACTC 867

RESULT 4
US-09-308-003-46/c
; Sequence 46, Application US/09308003
; Patent No. 6326170
; GENERAL INFORMATION:
; APPLICANT: Burnham, Martin K. R.
; APPLICANT: Lonetto, Michael A.
; APPLICANT: Warren, Patrick V.
; TITLE OF INVENTION: NOVEL PROKARYOTIC POLYNUCLEOTIDES,
; FILE REFERENCE: POLYPEPTIDES AND THEIR USES
; FILE REFERENCE: GM10093
; CURRENT APPLICATION NUMBER: US/09/308,003
; CURRENT FILING DATE: 1999-05-10
; EARLIER APPLICATION NUMBER: 60/058,710
; EARLIER FILING DATE: 1997-09-12
; NUMBER OF SEQ ID NOS: 52
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 46
; LENGTH: 933
; TYPE: DNA
; ORGANISM: Staphylococcus aureus
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (1)...(933)
US-09-308-003-46

Alignment Scores:
Pred. No.: 50 Length: 933
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x US-09-308-003-46 (1-933)

Qy 4 HisSerValLeuSerPheLeu 10
Db 891 CANTCTGTTCTCAGCTTCTTG 871

RESULT 5
US-08-221-817-21
; Sequence 21, Application US/08221817
; Patent No. 5532151
; GENERAL INFORMATION:
; APPLICANT: Chantry, David
; APPLICANT: Gray, Patrick W.
; APPLICANT: Hoekstra, Merle F.
; TITLE OF INVENTION: A No. 5532151el G Protein-Coupled Receptor
; FILE REFERENCE: KINASE GRK6
; NUMBER OF SEQUENCES: 24
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; ADDRESSEE: Borun

```

```

; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/221.817
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/123,932
; FILING DATE: 17 SEP 1993
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 5532151and, Greta E.
; REGISTRATION NUMBER: 35,302
; REFERENCE/DOCKET NUMBER: 31981
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1983 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 13...1740
US-08-221-817-21

Alignment Scores:
Pred. No.: 103 Length: 1983
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 1 Gaps: 0

US-10-799-747-116 (1-20) x US-08-221-817-21 (1-1983)

Qy 3 AlaHisSerValLeuSerPhe 9
Db 1262 GCTCACTCTGTTCTCAGCTTC 1282

RESULT 6
US-08-454-439-21
; Sequence 21, Application US/08454439
; Patent No. 5591618
; GENERAL INFORMATION:
; APPLICANT: Chantry, David
; APPLICANT: Gray, Patrick W.
; APPLICANT: Hoekstra, Merle F.
; TITLE OF INVENTION: A No. 5591618el G Protein-Coupled Receptor
; FILE REFERENCE: KINASE GRK6
; NUMBER OF SEQUENCES: 24
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; ADDRESSEE: Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM: disk
; MEDIUM TYPE: Floppy
; COMPUTER: IBM PC compatible

```

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; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/454,439
; FILING DATE: 30-MAY-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/221,817
; FILING DATE: 31-MAR-1994
; APPLICATION NUMBER: 08/123,932
; FILING DATE: 17 SEP 1993
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 5591618and, Greta E.
; REGISTRATION NUMBER: 35,302
; REFERENCE/DOCKET NUMBER: 31981
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1983 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 13..1740
US-08-454-439-21

Alignment Scores:
Pred. No.: 103 Length: 1983
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 1 Gaps: 0

US-10-799-747-116 (1-20) x US-08-454-439-21 (1-1983)
QY 3 AlaHisSerValLeuSerPhe 9
Db 1262 GCTCACTCTGTTCTCAGCTTC 1282

RESULT 7
PCT-US94-10487-21
; Sequence 21, Application PC/TUS9410487
; GENERAL INFORMATION:
; APPLICANT: ICOS Corporation
; TITLE OF INVENTION: A Novel G Protein-Coupled Receptor
; TITLE OF INVENTION: Kinase GRK6
; NUMBER OF SEQUENCES: 24
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Marshall, O'Toole, Gerstein, Murray &
; ADDRESSEE: Borun
; STREET: 6300 Sears Tower, 233 South Wacker Drive
; CITY: Chicago
; STATE: Illinois
; COUNTRY: USA
; ZIP: 60606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/10487
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/221,817

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; FILING DATE: 31 MAR 1994
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/123,932
; FILING DATE: 17 SEP 1993
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Noland, Greta E.
; REGISTRATION NUMBER: 35,302
; REFERENCE/DOCKET NUMBER: 27866/31981
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (312) 474-6300
; TELEFAX: (312) 474-0448
; TELEX: 25-3856
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1983 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 13..1740
PCT-US94-10487-21

Alignment Scores:
Pred. No.: 103 Length: 1983
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 5 Gaps: 0

US-10-799-747-116 (1-20) x PCT-US94-10487-21 (1-1983)
QY 3 AlaHisSerValLeuSerPhe 9
Db 1262 GCTCACTCTGTTCTCAGCTTC 1282

RESULT 8
US-08-836-442-1
; Sequence 1, Application US/08836442
; Patent No. 5990293
; GENERAL INFORMATION:
; APPLICANT: DOCHERTY, Andrew, J.P.
; APPLICANT: SLOCOMBE, Patrick, M.
; TITLE OF INVENTION: A HUMAN METALLOPROTEINASE
; TITLE OF INVENTION: VARIANTS THEREOF AND DNA SEQUENCES CODING THEREFOR
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: DIXE, BRONSTEIN, ROBERTS & CUSMAN, LLP
; STREET: 130 Water Street
; CITY: Boston
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,442
; FILING DATE: 01-MAY-1997
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/GB96/02181
; FILING DATE: 13-MAR-1997
; APPLICATION NUMBER: GB 9612150.4
; FILING DATE: 11-JUN-1996
; APPLICATION NUMBER: GB 9526229.1
; FILING DATE: 21-DEC-1995

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; APPLICATION NUMBER: GB 9521498.7
; FILING DATE: 20-OCT-1995
; APPLICATION NUMBER: GB 95521495.3
; FILING DATE: 20-OCT-1995
; APPLICATION NUMBER: GB 9518023.8
; FILING DATE: 05-SEP-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Resnick, David S
; REGISTRATION NUMBER: 34,235
; REFERENCE/DOCKET NUMBER: 47425
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-523-3400
; TELEFAX: 617-523-6440
; TELEX:
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2056 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-836-442-1
Alignment Scores:
Pred. No.: 107 Length: 2056
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 2 Gaps: 0
US-10-799-747-116 (1-20) x US-08-836-442-1 (1-2056)
QY 4 HisSerValneuSerPheLeu 10
Db 1656 CATTCGTCTATCCTTCTTA 1676
RESULT 9
US-09-786-256C-29
; Sequence 29, Application US/09786256C
; Patent No. 6680189
; GENERAL INFORMATION:
; APPLICANT: YOSHIMURA, Koji
; APPLICANT: HIKICHI, Yuichi
; APPLICANT: NISHIMURA, Atsushi
; TITLE OF INVENTION: No. 6680189el Protein and DNA Thereof
; FILE REFERENCE: 2544 USOP
; CURRENT APPLICATION NUMBER: US/09/786,256C
; PRIOR FILING DATE: 2001-03-02
; PRIOR APPLICATION NUMBER: PCT/JP99/04766
; PRIOR FILING DATE: 1999-09-02
; PRIOR APPLICATION NUMBER: JP 10-250115
; PRIOR FILING DATE: 1998-09-03
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 29
; LENGTH: 2560
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (55)..(1674)
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(2560)
; OTHER INFORMATION: DNA sequence of FIG 1-2 containing SEQ ID NO:3 encoding for prote
; FEATURE:
; NAME/KEY: mat_peptide
; LOCATION: (55)..()
US-09-786-256C-29
Alignment Scores:
Pred. No.: 131 Length: 2560
Score: 7.00 Matches: 2569
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0
US-10-799-747-116 (1-20) x US-08-956-171E-74 (1-2569)
QY 4 HisSerValneuSerPheLeu 10
Db 1703 CATTCGTCTATCCTTCTTA 1723
RESULT 10
US-08-956-171E-74
; Sequence 74, Application US/08956171E
; Patent No. 6593114
; GENERAL INFORMATION:
; APPLICANT: Charles Kunsch
; Gil H. Choi
; Patrick S. Dillon
; Craig A. Rosen
; Steven C. Barash
; Michael R. Fannon
; TITLE OF INVENTION: Staphylococcus aureus Polynucleotides and Sequences
; NUMBER OF SEQUENCES: 5256
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Human Genome Sciences, Inc.
; STREET: 9410 Key West Avenue
; CITY: Rockville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20850
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette, 3.50 inch, 1.4mb storage
; COMPUTER: HP Vectra 486/33
; OPERATING SYSTEM: MSDOS version 6.2
; SOFTWARE: ASCII Text
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/956,171E
; FILING DATE: 20-Oct-1997
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 60/009,861
; FILING DATE: January 5, 1996
; APPLICATION NUMBER: 08/781,986
; FILING DATE: January 3, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Mark J. Hyman
; REGISTRATION NUMBER: 46,789
; REFERENCE/DOCKET NUMBER: PH248P1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (240) 314-1224
; TELEFAX: (301) 309-8439
; INFORMATION FOR SEQ ID NO: 74:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2569 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 74:
US-08-956-171E-74
Alignment Scores:
Pred. No.: 132 Length: 2569
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0
US-10-799-747-116 (1-20) x US-08-956-171E-74 (1-2569)
QY 4 HisSerValneuSerPheLeu 10
```

```
Db 1609 CATTCTGTTCTCAGCTTCTTG 1629
|||||
RESULT 11
US-08-836-943-1/c
; Sequence 1, Application US/08836943
; Patent No. 5965391
; GENERAL INFORMATION:
; APPLICANT: Reinscheid, Dieter
; APPLICANT: Eikmanns, Bernhard
; APPLICANT: Sahm, Hermann
; TITLE OF INVENTION: DNA WHICH REGULATES GENE EXPRESSION IN
; CORYNEFORM BACTERIA
; NUMBER OF SEQUENCES: 3
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: The Firm of Karl F. Ross, PC
; STREET: 5676 Riverdale Ave.
; CITY: Bronx
; STATE: New York
; COUNTRY: USA
; ZIP: 10471
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/836,943
; FILING DATE: 08-MAY-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Myers, Jonathan
; REGISTRATION NUMBER: 26,963
; REFERENCE/DOCKET NUMBER: 20357
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (718) 884-6600
; TELEFAX: 718/601-1099
; TELEX: 620428
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3024 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Corynebacterium glutamicum
; STRAIN: ATCC 13032
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 586..2805
; US-08-836-943-1

Alignment Scores:
Pred. No.: 154 Length: 3024
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 2 Gaps: 0

US-10-799-747-116 (1-20) x US-08-836-943-1 (1-3024)
Qy 5 SerValLeuSerPheLeuLeu 11
|||||
Db 2812 AGCGTGCTTAGTTTGTCTT 2792
|||||
RESULT 12
US-09-134-000C-2717/c
; Sequence 2717, Application US/09134000C
; Patent No. 6617156
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```
; GENERAL INFORMATION:
; APPLICANT: Lynn Doucette-Stamm et al
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
; FILE REFERENCE: 032796-032
; CURRENT APPLICATION NUMBER: US/09/134,000C
; CURRENT FILING DATE: 1998-08-13
; PRIOR APPLICATION NUMBER: US 60/055,778
; PRIOR FILING DATE: 1997-08-15
; NUMBER OF SEQ ID NOS: 6812
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 2717
; LENGTH: 3465
; TYPE: DNA
; ORGANISM: Enterococcus faecalis
; US-09-134-000C-2717

Alignment Scores:
Pred. No.: 176 Length: 3465
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x US-09-134-000C-2717 (1-3465)
Qy 5 SerValLeuSerPheLeuLeu 11
|||||
Db 32 TCTGTTCTCTCTCTTTT 12
|||||
RESULT 13
US-08-569-214-1/c
; Sequence 1, Application US/08569214
; Patent No. 6165469
; GENERAL INFORMATION:
; APPLICANT: MANN, BARBARA J.
; APPLICANT: PETRI, WILLIAM A.
; TITLE OF INVENTION: RECOMBINANT ENTAMOEBA HISTOLYTICA LECTIN
; TITLE OF INVENTION: SUBUNIT PEPTIDES AND REAGENTS SPECIFIC FOR MEMBERS OF THE
; TITLE OF INVENTION: 170 KD SUBUNIT MULTIGENE FAMILY
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 2000 PENNSYLVANIA AVENUE N.W., STE. 5500
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: USA
; ZIP: 20006-1812
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/569,214
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US94/06890
; FILING DATE: 17-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: MURASHIGE, KATE H.
; REGISTRATION NUMBER: 29,959
; REFERENCE/DOCKET NUMBER: 9148-0006.21
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 887-1500
; TELEFAX: (202) 887-0763
; TELEX: 90-4030
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3892 base pairs
; TYPE: nucleic acid
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```

; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: CDS
; LOCATION: join(1..3873, 3877..3882, 3886..3891)
US-08-569-214-1
Alignment Scores:
Pred. No.: 196 Length: 3892
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-569-214-1 (1-3892)

QY 4 HisSerValLeuSerPheLeu 10
Db 230 CATTCGTATTATCATTTTITA 210

RESULT 14
US-08-937-236-1/c
; Sequence 1, Application US/08937236
; Patent No. 6187310
; GENERAL INFORMATION:
; APPLICANT: MANN, BARBARA J.
; APPLICANT: PETRI, WILLIAM A.
; APPLICANT: DODSON, JAMES M.
; TITLE OF INVENTION: RECOMBINANT ENTAMORIBA HISTOLYTICA LECTIN
; TITLE OF INVENTION: SUBUNIT PEPTIDES AND REAGENTS SPECIFIC FOR MEMBERS OF THE
; TITLE OF INVENTION: 170 KD SUBUNIT MULTIGENE FAMILY
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 2000 PENNSYLVANIA AVENUE N.W., STE. 5500
; CITY: WASHINGTON
; STATE: DC
; COUNTRY: USA
; ZIP: 20006-1812
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/937,236
; FILING DATE:
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/569,214
; FILING DATE: 16 SEPTEMBER 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: LIVNAT, SHMUEL
; REGISTRATION NUMBER: 33,949
; REFERENCE/DOCKET NUMBER: 291482000622
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (202) 887-1500
; TELEFAX: (202) 887-0763
; TELEX: 90-4030
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 3892 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; FEATURE:
; NAME/KEY: CDS
; LOCATION: join(1..3873, 3877..3882, 3886..3891)
US-08-937-236-1
Alignment Scores:
Pred. No.: 196 Length: 3892

```

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Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 3 Gaps: 0

US-10-799-747-116 (1-20) x US-08-937-236-1 (1-3892)

QY 4 HisSerValLeuSerPheLeu 10
Db 230 CATTCGTATTATCATTTTITA 210

RESULT 15
US-09-081-385-8/c
; Sequence 8, Application US/09081385
; Patent No. 6593456
; GENERAL INFORMATION:
; APPLICANT: Gatanaga, T.
; APPLICANT: Granger, G.A.
; TITLE OF INVENTION: Factors Altering Tumor Necrosis
; TITLE OF INVENTION: Factor Receptor Releasing Enzyme Activity, and Methods
; TITLE OF INVENTION: of Use Thereof
; NUMBER OF SEQUENCES: 154
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: MORRISON & FOERSTER
; STREET: 755 PAGE MILL ROAD
; CITY: Palo Alto
; STATE: CA
; COUNTRY: USA
; ZIP: 94304-1018
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/081,385
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/964,747
; FILING DATE: 05-NOV-1997
; APPLICATION NUMBER: 60/030,761
; FILING DATE: 06-NOV-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Wu, Frank
; REGISTRATION NUMBER: 41,396
; REFERENCE/DOCKET NUMBER: 22000-20577.21
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-813-5600
; TELEFAX: 650-494-0792
; TELEX: 706141
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 4218 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: Genomic DNA
US-09-081-385-8
Alignment Scores:
Pred. No.: 212 Length: 4218
Score: 7.00 Matches: 7
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 35.00% Indels: 0
DB: 4 Gaps: 0

US-10-799-747-116 (1-20) x US-09-081-385-8 (1-4218)

QY 5 SerValLeuSerPheLeu 11

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Db 449 TCAGTGTATCCTTCCTTA 429

Search completed: July 21, 2004, 05:24:13  
Job time : 94 secs

GenCore version 5.1.6  
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OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 04:43:08 ; Search time 3357 Seconds  
(without alignments)

29.065 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 20

Sequence: 1 MAHVSLSFLWTYPYALKSX 20

Scoring table:

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Ygapop 60.0 , Ygapext 60.0  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 3191023 seqs, 2439312756 residues

Word size: 1

Total number of hits satisfying chosen parameters: 6371784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

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-TRANS=human40.cdi -LIST=45 -DOALIGN=200 -THR SCORE=quality -THR MIN=1  
-ALIGN=15 -MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZ=500 -MINLEN=0  
-MAXLEN=2000000000 -USER=US10799747@CGN 1 1 723 @runat 19072004 161446 22092  
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-FGAPOP=6 -FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

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Published Applications NA:  
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7: /cgn2\_6/ptodata/1/pubpna/US08\_NEW\_PUB.seq.\*  
8: /cgn2\_6/ptodata/1/pubpna/US08\_PUBCOMB.seq.\*  
9: /cgn2\_6/ptodata/1/pubpna/US09A\_PUBCOMB.seq.\*  
10: /cgn2\_6/ptodata/1/pubpna/US09B\_PUBCOMB.seq.\*  
11: /cgn2\_6/ptodata/1/pubpna/US09C\_PUBCOMB.seq.\*  
12: /cgn2\_6/ptodata/1/pubpna/US09\_NEW\_PUB.seq.\*  
13: /cgn2\_6/ptodata/1/pubpna/US09\_NEW\_PUB.seq2.\*  
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16: /cgn2\_6/ptodata/1/pubpna/US10C\_PUBCOMB.seq.\*  
17: /cgn2\_6/ptodata/1/pubpna/US10D\_PUBCOMB.seq.\*  
18: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq.\*  
19: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq2.\*  
20: /cgn2\_6/ptodata/1/pubpna/US10\_NEW\_PUB.seq3.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
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1	19	95.0	1434	15	US-10-195-730-12	Sequence 12, Appl
2	8	40.0	356	9	US-09-867-701-768	Sequence 768, App
C 3	8	40.0	670	13	US-10-027-632-187894	Sequence 187894,
C 4	8	40.0	670	16	US-10-027-632-187894	Sequence 187894,
C 5	8	40.0	740	13	US-10-027-632-11891	Sequence 11891, A
C 6	8	40.0	740	16	US-10-027-632-11891	Sequence 11891, A
7	8	40.0	1627	17	US-10-437-963-82948	Sequence 82948, A
8	8	40.0	2037	17	US-10-437-963-82948	Sequence 82948, A
9	8	40.0	6681	17	US-10-322-281-411	Sequence 411, App
C 10	8	40.0	133787	17	US-10-322-281-411	Sequence 858, App
C 11	8	40.0	196063	17	US-10-322-281-612	Sequence 612, App
C 12	8	40.0	1601042	13	US-10-027-632-59064	Sequence 59064, A
C 13	8	40.0	1601042	16	US-10-027-632-59064	Sequence 59064, A
14	7	35.0	65	15	US-10-032-585-3022	Sequence 3022, Ap
15	7	35.0	163	13	US-10-424-599-127928	Sequence 127928,
C 16	7	35.0	191	13	US-10-424-599-98581	Sequence 98581, A
17	7	35.0	221	13	US-10-085-783A-36161	Sequence 36161, A
C 18	7	35.0	221	16	US-10-242-535A-36161	Sequence 36161, A
19	7	35.0	240	13	US-10-424-599-132181	Sequence 132181,
C 20	7	35.0	279	13	US-10-424-599-22323	Sequence 22323, A
C 21	7	35.0	292	13	US-10-424-599-133734	Sequence 133734,
C 22	7	35.0	293	11	US-09-864-408A-2007	Sequence 2007, Ap
C 23	7	35.0	315	14	US-10-040-739-894	Sequence 894, App
24	7	35.0	324	13	US-10-027-632-143650	Sequence 143650,
25	7	35.0	324	16	US-10-027-632-143650	Sequence 143650,
26	7	35.0	348	17	US-10-437-963-77562	Sequence 77562, A
C 27	7	35.0	352	9	US-09-770-791-678	Sequence 678, App
C 28	7	35.0	379	10	US-09-918-995-37721	Sequence 37721, A
C 29	7	35.0	415	17	US-10-437-963-38874	Sequence 38874, A
C 30	7	35.0	423	13	US-10-027-632-104598	Sequence 104598,
31	7	35.0	423	13	US-10-027-632-298016	Sequence 298016,
C 32	7	35.0	423	16	US-10-027-632-104598	Sequence 104598,
C 33	7	35.0	423	16	US-10-027-632-298016	Sequence 298016,
C 34	7	35.0	431	17	US-10-437-963-44015	Sequence 44015, A
C 35	7	35.0	432	9	US-09-974-300-3604	Sequence 3604, Ap
C 36	7	35.0	433	13	US-09-770-423-79	Sequence 79, Appl
C 37	7	35.0	435	13	US-10-424-599-41140	Sequence 41140, A
C 38	7	35.0	440	10	US-09-918-995-19984	Sequence 19984, A
C 39	7	35.0	442	13	US-10-424-599-59524	Sequence 59524, A
C 40	7	35.0	445	10	US-09-918-995-29596	Sequence 29596, A
C 41	7	35.0	455	13	US-09-918-995-2636	Sequence 2636, Ap
42	7	35.0	455	13	US-10-027-632-179616	Sequence 179616,
43	7	35.0	455	16	US-10-027-632-179616	Sequence 179616,
44	7	35.0	465	9	US-09-864-761-1120	Sequence 1120, Ap
C 45	7	35.0	465	9	US-09-796-692-7472	Sequence 7472, Ap

#### ALIGNMENTS

#### RESULT 1

US-10-195-730-12  
; Sequence 12, Application US/10195730  
; Publication No. US2003014492A1  
; GENERAL INFORMATION:  
; APPLICANT: Rosen et. al  
; TITLE OF INVENTION: 101 Human Secreted Proteins  
; FILE REFERENCE: P2017P1  
; CURRENT APPLICATION NUMBER: US/10/195,730  
; CURRENT FILING DATE: 2002-07-16  
; PRIOR FILING DATE: 1999-03-31  
; PRIOR APPLICATION NUMBER: 60/060,837  
; PRIOR FILING DATE: 1997-10-02  
; PRIOR APPLICATION NUMBER: 60/060,862  
; NUMBER OF SEQ ID NOS: 390  
; SOFTWARE: PatentIn Ver. 2.0  
; SEQ ID NO 12  
; LENGTH: 1434  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-10-195-730-12

Alignment Scores: 1.22e-10 1434  
Pred. No.: 19.00 Matches: 19  
Score: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 95.00% Indels: 0  
DB: 15 Gaps: 0

US-10-799-747-116 (1-20) x US-10-195-730-12 (1-1434)

Qy 1 MetAlaHisSerValLeuSerPheLeuLeuTrpThrProTyAlaLeuLysSer 19  
Db 507 ATGCAGCCCATTCAGTCTTGAGTTTCTTCTCGACACCTTATGCTCTGAATCA 563

## RESULT 2

US-09-867-701-768  
; Sequence 768, Application US/09867701  
; Patent No. US20020132237A1  
; GENERAL INFORMATION:  
; APPLICANT: Aglate, Paul A.  
; APPLICANT: Jones, Robert  
; APPLICANT: Harlocker, Susan L.  
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY  
; TITLE OF INVENTION: AND DIAGNOSIS OF OVARIAN CANCER  
; FILE REFERENCE: 210121.497  
; CURRENT APPLICATION NUMBER: US/09/867,701  
; CURRENT FILING DATE: 2001-05-29  
; NUMBER OF SEQ ID NOS: 10912  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 768  
; LENGTH: 356  
; TYPE: DNA  
; ORGANISM: Homo sapien  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: (1)...(356)  
; OTHER INFORMATION: n = A,T,C or G  
US-09-867-701-768

Alignment Scores: 11.2 Length: 356  
Pred. No.: 8.00 Matches: 8  
Score: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x US-09-867-701-768 (1-356)

Qy 3 AlaHisSerValLeuSerPheLeu 10  
Db 108 GTCATTCGTACTTCTCTCTC 131

## RESULT 3

US-10-027-632-187894/c  
; Sequence 187894, Application US/10027632  
; Publication No. US20020198371A1  
; GENERAL INFORMATION:  
; APPLICANT: Wang, David G.  
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
; TITLE OF INVENTION: Polymorphisms in the Human Genome  
; FILE REFERENCE: 108827.129  
; CURRENT APPLICATION NUMBER: US/10/027,632  
; CURRENT FILING DATE: 2002-04-30  
; PRIOR APPLICATION NUMBER: US 60/218,006  
; PRIOR FILING DATE: 2000-07-12  
; PRIOR APPLICATION NUMBER: US 60/198,676  
; PRIOR FILING DATE: 2000-04-20  
; PRIOR APPLICATION NUMBER: US 60/193,483  
; PRIOR FILING DATE: 2000-03-29  
; PRIOR APPLICATION NUMBER: US 60/185,218  
; PRIOR FILING DATE: 2000-02-24

; PRIOR APPLICATION NUMBER: US 60/167,363  
; PRIOR FILING DATE: 1999-11-23  
; PRIOR APPLICATION NUMBER: US 60/156,358  
; PRIOR FILING DATE: 1999-09-28  
; PRIOR APPLICATION NUMBER: US 60/146,002  
; PRIOR FILING DATE: 1999-08-09  
; NUMBER OF SEQ ID NOS: 325720  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 187894  
; LENGTH: 670  
; TYPE: DNA  
; ORGANISM: Human  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: (1)...(670)  
; OTHER INFORMATION: n = A,T,C or G  
US-10-027-632-187894

Alignment Scores: 20 Length: 670  
Pred. No.: 8.00 Matches: 8  
Score: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-187894 (1-670)

Qy 1 MetAlaHisSerValLeuSer 8  
Db 36 ATGCAGCCCATAGTGTGCTCTCC 13

## RESULT 4

US-10-027-632-187894/c  
; Sequence 187894, Application US/10027632  
; Publication No. US20030204075A9  
; GENERAL INFORMATION:  
; APPLICANT: Wang, David G.  
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
; TITLE OF INVENTION: Polymorphisms in the Human Genome  
; FILE REFERENCE: 108827.129  
; CURRENT APPLICATION NUMBER: US/10/027,632  
; CURRENT FILING DATE: 2002-04-30  
; PRIOR APPLICATION NUMBER: US 60/218,006  
; PRIOR FILING DATE: 2000-07-12  
; PRIOR APPLICATION NUMBER: US 60/198,676  
; PRIOR FILING DATE: 2000-04-20  
; PRIOR APPLICATION NUMBER: US 60/193,483  
; PRIOR FILING DATE: 2000-03-29  
; PRIOR APPLICATION NUMBER: US 60/185,218  
; PRIOR FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: US 60/167,363  
; PRIOR FILING DATE: 1999-11-23  
; PRIOR APPLICATION NUMBER: US 60/156,358  
; PRIOR FILING DATE: 1999-09-28  
; PRIOR APPLICATION NUMBER: US 60/146,002  
; PRIOR FILING DATE: 1999-08-09  
; NUMBER OF SEQ ID NOS: 325720  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 187894  
; LENGTH: 670  
; TYPE: DNA  
; ORGANISM: Human  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: (1)...(670)  
; OTHER INFORMATION: n = A,T,C or G  
US-10-027-632-187894

Alignment Scores: 20 Length: 670  
Pred. No.: 8.00 Matches: 8  
Score: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 40.00% Indels: 0  
 DB: 16 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-187894 (1-670)

Qy 1 MetAlaAlaHisSerValLeuSer 8  
 Db 36 ATGGCAGCCCATAGTGTCTCTCC 13

## RESULT 5

US-10-027-632-11891/c  
 ; Sequence 11891, Application US/10027632  
 ; Publication No. US20020198371A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Wang, David G.  
 ; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
 ; FILE REFERENCE: 108827.129  
 ; CURRENT APPLICATION NUMBER: US/10/027,632  
 ; CURRENT FILING DATE: 2002-04-30  
 ; PRIOR APPLICATION NUMBER: US 60/218,006  
 ; PRIOR FILING DATE: 2000-07-12  
 ; PRIOR APPLICATION NUMBER: US 60/198,676  
 ; PRIOR FILING DATE: 2000-04-20  
 ; PRIOR APPLICATION NUMBER: US 60/193,483  
 ; PRIOR FILING DATE: 2000-03-29  
 ; PRIOR APPLICATION NUMBER: US 60/185,218  
 ; PRIOR FILING DATE: 2000-02-24  
 ; PRIOR APPLICATION NUMBER: US 60/167,363  
 ; PRIOR FILING DATE: 1999-11-23  
 ; PRIOR APPLICATION NUMBER: US 60/156,358  
 ; PRIOR FILING DATE: 1999-09-28  
 ; PRIOR APPLICATION NUMBER: US 60/146,002  
 ; PRIOR FILING DATE: 1999-08-09  
 ; NUMBER OF SEQ ID NOS: 325720  
 ; SOFTWARE: FastSeq for Windows Version 4.0  
 ; SEQ ID NO 11891  
 ; LENGTH: 740  
 ; TYPE: DNA  
 ; ORGANISM: Human  
 ; US-10-027-632-11891

Alignment Scores:  
 Pred. No.: 21.9 Length: 740  
 Score: 8.00 Matches: 8  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 40.00% Indels: 0  
 DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-11891 (1-740)

Qy 5 SerValLeuSerPheLeuLeuTrp 12  
 Db 266 AGTGTCTTAAGCTTCCTCTCTGG 243

## RESULT 6

US-10-027-632-11891/c  
 ; Sequence 11891, Application US/10027632  
 ; Publication No. US20030204075A9  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Wang, David G.  
 ; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide  
 ; FILE REFERENCE: 108827.129  
 ; CURRENT APPLICATION NUMBER: US/10/027,632  
 ; CURRENT FILING DATE: 2002-04-30  
 ; PRIOR APPLICATION NUMBER: US 60/218,006  
 ; PRIOR FILING DATE: 2000-07-12  
 ; PRIOR APPLICATION NUMBER: US 60/198,676  
 ; PRIOR FILING DATE: 2000-04-20  
 ; PRIOR APPLICATION NUMBER: US 60/193,483

; PRIOR FILING DATE: 2000-03-29  
 ; PRIOR APPLICATION NUMBER: US 60/185,218  
 ; PRIOR FILING DATE: 2000-02-24  
 ; PRIOR APPLICATION NUMBER: US 60/167,363  
 ; PRIOR FILING DATE: 1999-11-23  
 ; PRIOR APPLICATION NUMBER: US 60/156,358  
 ; PRIOR FILING DATE: 1999-09-28  
 ; PRIOR APPLICATION NUMBER: US 60/146,002  
 ; PRIOR FILING DATE: 1999-08-09  
 ; NUMBER OF SEQ ID NOS: 325720  
 ; SOFTWARE: FastSeq for Windows Version 4.0  
 ; SEQ ID NO 11891  
 ; LENGTH: 740  
 ; TYPE: DNA  
 ; ORGANISM: Human  
 ; US-10-027-632-11891

Alignment Scores:  
 Pred. No.: 21.9 Length: 740  
 Score: 8.00 Matches: 8  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 40.00% Indels: 0  
 DB: 16 Gaps: 0

US-10-799-747-116 (1-20) x US-10-027-632-11891 (1-740)

Qy 5 SerValLeuSerPheLeuLeuTrp 12  
 Db 266 AGTGTCTTAAGCTTCCTCTCTGG 243

## RESULT 7

US-10-437-963-82948  
 ; Sequence 82948, Application US/10437963  
 ; Publication No. US20040123343A1  
 ; GENERAL INFORMATION:  
 ; APPLICANT: La Rosa, Thomas J.  
 ; APPLICANT: Kovalic, David K.  
 ; APPLICANT: Zhou, Yihua  
 ; APPLICANT: Cao, Yongwei  
 ; APPLICANT: Wu, Wei  
 ; APPLICANT: Boukharov, Andrey A.  
 ; APPLICANT: Barbazuk, Brad  
 ; APPLICANT: Li, Ping  
 ; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With  
 ; FILE REFERENCE: 38-21(53221)B  
 ; CURRENT APPLICATION NUMBER: US/10/437,963  
 ; CURRENT FILING DATE: 2003-05-14  
 ; NUMBER OF SEQ ID NOS: 204966  
 ; SEQ ID NO 82948  
 ; LENGTH: 1627  
 ; TYPE: DNA  
 ; ORGANISM: Oryza sativa  
 ; FEATURE:  
 ; OTHER INFORMATION: Clone ID: PAT\_MRT4530\_82327C.1  
 ; US-10-437-963-82948

Alignment Scores:  
 Pred. No.: 45.2 Length: 1627  
 Score: 8.00 Matches: 8  
 Percent Similarity: 100.00% Conservative: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 40.00% Indels: 0  
 DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-437-963-82948 (1-1627)

Qy 3 AlaHisSerValLeuSerPheLeu 10  
 Db 266 GCACACAGTGTTCCTCTCTT 289

## RESULT 8

```
US-10-437-963-82916
; Sequence 82916, Application US/10437963
; Publication No. US20040123343A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; APPLICANT: Wu, Wei
; APPLICANT: Boukharov, Andrey A.
; APPLICANT: Barabazuk, Brad
; APPLICANT: Li, Ping
; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/437,963
; CURRENT FILING DATE: 2003-05-14
; NUMBER OF SEQ ID NOS: 204966
; SEQ ID NO 82916
; LENGTH: 2037
; TYPE: DNA
; ORGANISM: Oryza sativa
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MKT4530_82299C.1
US-10-437-963-82916

Alignment Scores:
Pred. No.: 55.6 Length: 2037
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-437-963-82916 (1-2037)

QY 6 ValLeuSerPheLeuThr 13
Db 69 GTGCTATCCTTCTCTCTGGACA 92

RESULT 9
US-10-322-281-411
; Sequence 411, Application US/10322281
; Publication No. US20040126762A1
; GENERAL INFORMATION:
; APPLICANT: David W. Morris
; APPLICANT: Marc S. Malandro
; TITLE OF INVENTION: Novel Compositions and Methods in Cancer
; FILE REFERENCE: 529452001000
; CURRENT APPLICATION NUMBER: US/10/322,281
; CURRENT FILING DATE: 2002-12-17
; NUMBER OF SEQ ID NOS: 866
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 411
; LENGTH: 66681
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(66681)
; OTHER INFORMATION: n = A,T,C or G
US-10-322-281-411

Alignment Scores:
Pred. No.: 1.37e+03 Length: 66681
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-322-281-411 (1-66681)

US-10-437-963-82916
; Sequence 82916, Application US/10437963
; Publication No. US20040123343A1
; GENERAL INFORMATION:
; APPLICANT: La Rosa, Thomas J.
; APPLICANT: Kovalic, David K.
; APPLICANT: Zhou, Yihua
; APPLICANT: Cao, Yongwei
; APPLICANT: Wu, Wei
; APPLICANT: Boukharov, Andrey A.
; APPLICANT: Barabazuk, Brad
; APPLICANT: Li, Ping
; TITLE OF INVENTION: Rice Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53221)B
; CURRENT APPLICATION NUMBER: US/10/437,963
; CURRENT FILING DATE: 2003-05-14
; NUMBER OF SEQ ID NOS: 204966
; SEQ ID NO 82916
; LENGTH: 2037
; TYPE: DNA
; ORGANISM: Oryza sativa
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MKT4530_82299C.1
US-10-437-963-82916

Alignment Scores:
Pred. No.: 55.6 Length: 2037
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-437-963-82916 (1-2037)

QY 6 ValLeuSerPheLeuThr 13
Db 69 GTGCTATCCTTCTCTCTGGACA 92

RESULT 9
US-10-322-281-411
; Sequence 411, Application US/10322281
; Publication No. US20040126762A1
; GENERAL INFORMATION:
; APPLICANT: David W. Morris
; APPLICANT: Marc S. Malandro
; TITLE OF INVENTION: Novel Compositions and Methods in Cancer
; FILE REFERENCE: 529452001000
; CURRENT APPLICATION NUMBER: US/10/322,281
; CURRENT FILING DATE: 2002-12-17
; NUMBER OF SEQ ID NOS: 866
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 411
; LENGTH: 66681
; TYPE: DNA
; ORGANISM: Mus musculus
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(66681)
; OTHER INFORMATION: n = A,T,C or G
US-10-322-281-411

Alignment Scores:
Pred. No.: 1.37e+03 Length: 66681
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-322-281-411 (1-66681)

QY 4 HisSerValLeuSerPheLeu 11
Db 38196 CACTCTGTCTTCATTCCTTTG 38219

RESULT 10
US-10-322-281-858/c
; Sequence 858, Application US/10322281
; Publication No. US20040126762A1
; GENERAL INFORMATION:
; APPLICANT: David W. Morris
; APPLICANT: Marc S. Malandro
; TITLE OF INVENTION: Novel Compositions and Methods in Cancer
; FILE REFERENCE: 529452001000
; CURRENT APPLICATION NUMBER: US/10/322,281
; CURRENT FILING DATE: 2002-12-17
; NUMBER OF SEQ ID NOS: 866
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 858
; LENGTH: 133787
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(133787)
; OTHER INFORMATION: n = A,T,C or G
US-10-322-281-858

Alignment Scores:
Pred. No.: 2.61e+03 Length: 133787
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-322-281-858 (1-133787)

QY 4 HisSerValLeuSerPheLeu 11
Db 35487 CACTCAGTTCATCATCTTCCTT 35464

RESULT 11
US-10-322-281-612
; Sequence 612, Application US/10322281
; Publication No. US20040126762A1
; GENERAL INFORMATION:
; APPLICANT: David W. Morris
; APPLICANT: Marc S. Malandro
; TITLE OF INVENTION: Novel Compositions and Methods in Cancer
; FILE REFERENCE: 529452001000
; CURRENT APPLICATION NUMBER: US/10/322,281
; CURRENT FILING DATE: 2002-12-17
; NUMBER OF SEQ ID NOS: 866
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 612
; LENGTH: 196063
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)...(196063)
; OTHER INFORMATION: n = A,T,C or G
US-10-322-281-612

Alignment Scores:
Pred. No.: 3.7e+03 Length: 196063
Score: 8.00 Matches: 8
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 40.00% Indels: 0
DB: 17 Gaps: 0

US-10-799-747-116 (1-20) x US-10-322-281-612 (1-196063)
```

```

QY      3 AlahisServalLeuSerPheLeu 10
Db      50669 GCTCATTCTGTACTTTCCTTCCTC 50692

RESULT 12
US-10-027-632-59064/c
; Sequence 59064, Application US/10027632
; Publication No. US20020198371A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-04-20
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29
; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 59064
; LENGTH: 1601042
; TYPE: DNA
; ORGANISM: Human
; NAME/KEY: misc feature
; LOCATION: (1)...(1601042)
; OTHER INFORMATION: n = A,T,C or G
US-10-027-632-59064

Alignment Scores:
Pred. No.:      2.55e+04      Length:      1601042
Score:          8.00         Matches:      8
Percent Similarity: 100.00%   Conservative: 0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    40.00%       Indels:      0
DB:             13          Gaps:        0

US-10-799-747-116 (1-20) x US-10-027-632-59064 (1-1601042)

QY      3 AlahisServalLeuSerPheLeu 10
Db      638163 GCCCACTCGGTCTTAGCTTCTG 638140

RESULT 13
US-10-027-632-59064/c
; Sequence 59064, Application US/10027632
; Publication No. US20030204075A9
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
; TITLE OF INVENTION: Identification and Mapping of Single Nucleotide
; Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.129
; CURRENT APPLICATION NUMBER: US/10/027,632
; PRIOR APPLICATION NUMBER: US 60/218,006
; PRIOR FILING DATE: 2002-04-30
; PRIOR APPLICATION NUMBER: US 60/198,676
; PRIOR FILING DATE: 2000-07-12
; PRIOR APPLICATION NUMBER: US 60/193,483
; PRIOR FILING DATE: 2000-03-29

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; PRIOR APPLICATION NUMBER: US 60/185,218
; PRIOR FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/167,363
; PRIOR FILING DATE: 1999-11-23
; PRIOR APPLICATION NUMBER: US 60/156,358
; PRIOR FILING DATE: 1999-09-28
; PRIOR APPLICATION NUMBER: US 60/146,002
; PRIOR FILING DATE: 1999-08-09
; NUMBER OF SEQ ID NOS: 325720
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 59064
; LENGTH: 1601042
; TYPE: DNA
; ORGANISM: Human
; NAME/KEY: misc feature
; LOCATION: (1)...(1601042)
; OTHER INFORMATION: n = A,T,C or G
US-10-027-632-59064

Alignment Scores:
Pred. No.:      2.55e+04      Length:      1601042
Score:          8.00         Matches:      8
Percent Similarity: 100.00%   Conservative: 0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    40.00%       Indels:      0
DB:             16          Gaps:        0

US-10-799-747-116 (1-20) x US-10-027-632-59064 (1-1601042)

QY      3 AlahisServalLeuSerPheLeu 10
Db      638163 GCCCACTCGGTCTTAGCTTCTG 638140

RESULT 14
US-10-032-585-3022
; Sequence 3022, Application US/10032585
; Publication No. US20030180953A1
; GENERAL INFORMATION:
; APPLICANT: Terry, Roemer D.
; APPLICANT: Bo, Jiang
; APPLICANT: Charles, Boone
; APPLICANT: Howard, Bussey
; FILE OF INVENTION: Gene Disruption Methodologies for Drug Target Discovery
; FILE REFERENCE: 10182-005-999
; CURRENT APPLICATION NUMBER: US/10/032,585
; CURRENT FILING DATE: 2001-12-20
; NUMBER OF SEQ ID NOS: 8000
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 3022
; LENGTH: 65
; TYPE: DNA
; ORGANISM: Candida albicans
US-10-032-585-3022

Alignment Scores:
Pred. No.:      26.1         Length:      65
Score:          7.00         Matches:      7
Percent Similarity: 100.00%   Conservative: 0
Best Local Similarity: 100.00% Mismatches:    0
Query Match:    35.00%       Indels:      0
DB:             15          Gaps:        0

US-10-799-747-116 (1-20) x US-10-032-585-3022 (1-65)

QY      5 ServalLeuSerPheLeu 11
Db      13 TCTGTATTATCATCTTATTG 33

RESULT 15
US-10-424-599-127928
; Sequence 127928, Application US/10424599
; Publication No. US20040031072A1

```

```

; GENERAL INFORMATION:
; APPLICANT: La Rosa Thomas J
; APPLICANT: Kovalic David K
; APPLICANT: Zhou Yihua
; APPLICANT: Cao Yongwei
; TITLE OF INVENTION: Soy Nucleic Acid Molecules and Other Molecules Associated With
; TITLE OF INVENTION: Plants and Uses Thereof for Plant Improvement
; FILE REFERENCE: 38-21(53223)B
; CURRENT APPLICATION NUMBER: US/10/424,599
; CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 285684
; SEQ ID NO 127928
; LENGTH: 163
; TYPE: DNA
; ORGANISM: Glycine max
; FEATURE:
; OTHER INFORMATION: Clone ID: PAT_MRT3847_86523C.1
US-10-424-599-127928

Alignment Scores:
Pred. No.:      60.8      Length:      163
Score:          7.00      Matches:      7
Percent Similarity: 100.00%      Conservative: 0
Best Local Similarity: 100.00%      Mismatches: 0
Query Match:      35.00%      Indels:      0
DB:              13          Gaps:      0

US-10-799-747-116 (1-20) x US-10-424-599-127928 (1-163)

QY      4 HisSerValLeuSerPheLeu 10
      |||||
DB      40 CACTCTGTGTGTCCTTTCTG 60

```

Search completed: July 21, 2004, 07:54:36  
Job time : 3580 secs



GenCore version 5.1.6  
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM protein - nucleic search, using frame\_plus\_p2n model

Run on: July 21, 2004, 03:49:02 ; Search time 2467 Seconds  
(without alignments)  
242.093 Million cell updates/sec

Title: US-10-799-747-116

Perfect score: 20  
Sequence: 1 MAHSVLSFLWTFYALKSX 20

Scoring table:

OLIGO  
Xgapop 60.0 , Xgapext 60.0  
Ygapop 60.0 , Ygapext 60.0  
Fgapop 6.0 , Fgapext 7.0  
Delop 6.0 , Delext 7.0

Searched: 27513289 seqs, 14931090276 residues

Word size: 1

Total number of hits satisfying chosen parameters: 55023894

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Command line parameters:

-MODEL=frame+p2n.model -DEV=xlp  
-Q=/cgm2\_1/USPTO.spool\_p/US10799747/runat\_19072004.161443.21994/app\_query.fasta\_1.199  
-DB=EST -QPMT=fastcap -SUFFIX=olig.rst -MINMATCH=0.1 -LOOPCL=0 -LOOPEXT=0  
-UNITS=bits -START=1 -END=-1 -MATRIX=oligo -TRANS=human40.cdi -LIST=45  
-DOCALIGN=200 -THR SCORE=quality -THR\_MIN=1 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc  
-NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000  
-USER=US10799747@cgm2\_1.5180 @runat\_19072004.161443.21994 -NCPU=6 -ICPU=3  
-NO MMAP -LARGEQUERY -NEG SCORES=0 -WAIT -DSPBLOCK=100 -LONGLOG  
-DEV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=60 -XGAPEXT=60 -FGAPOP=6  
-FGAPEXT=7 -YGAPOP=60 -YGAPEXT=60 -DELOP=6 -DELEXT=7

Database :

EST:  
1: em\_estba:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estmu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_htc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_htc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: em\_gss\_hum:\*  
18: em\_gss\_inv:\*  
19: em\_gss\_pln:\*  
20: em\_gss\_vrt:\*  
21: em\_gss\_fun:\*  
22: em\_gss\_mam:\*  
23: em\_gss\_mus:\*  
24: em\_gss\_pro:\*  
25: em\_gss\_rod:\*  
26: em\_gss\_phg:\*  
27: em\_gss\_vrl:\*  
28: gb\_gss1:\*

29: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
C 1	19	95.0	240	10	BF910533	BF910533 CM4-UT004
2	19	95.0	575	12	BM722991	BM722991 UI-B-E01-
3	19	95.0	765	12	BI914473	BI914473 603182264
4	19	95.0	785	12	BG484396	BG484396 602505037
5	19	95.0	1201	13	BX441923	BX441923 BX441923
6	10	50.0	621	9	AL035941	AL035941 DKF2p564B
7	9	45.0	496	28	AQ293004	AQ293004 HS 2225 A
8	8	40.0	239	9	AV851019	AV851019 AV851019-
9	8	40.0	272	12	BG371993	BG371993 UI-R-CV0-
10	8	40.0	290	29	CE488576	CE488576 tigr-ges-
11	8	40.0	323	28	AQ350287	AQ350287 RPI11-13
12	8	40.0	328	29	CC808823	CC808823 ZMMB0047
13	8	40.0	336	14	CD202965	CD202965 MS1-0139P
14	8	40.0	356	14	T59368	T59368 yb57h01.s1
15	8	40.0	385	29	CE286752	CE286752 tigr-ges-
16	8	40.0	396	28	BH363737	BH363737 CH230-127
17	8	40.0	400	28	AQ475321	AQ475321 CITBI-E1-
18	8	40.0	419	13	BM204790	BM204790 BM204790
19	8	40.0	430	14	CD201054	CD201054 MS1-0124U
20	8	40.0	430	14	CD201131	CD201131 MS1-0124U
21	8	40.0	441	9	AV888988	AV888988 AV888988
22	8	40.0	444	10	BF412147	BF412147 UI-R-BF1-
23	8	40.0	448	14	CD113088	CD113088 ME1-0028T
24	8	40.0	448	14	CD201074	CD201074 MS1-0124U
25	8	40.0	448	14	CD201080	CD201080 MS1-0124U
26	8	40.0	448	14	CD201099	CD201099 MS1-0124U
27	8	40.0	448	14	CD201120	CD201120 MS1-0124U
C 28	8	40.0	457	9	AV981198	AV981198 AV981198
C 29	8	40.0	458	28	BZ615844	BZ615844 1955008.b
30	8	40.0	468	28	AQ945119	AQ945119 Sheared D
31	8	40.0	470	28	AQ702871	AQ702871 HS 5452 B
C 32	8	40.0	481	13	BM196597	BM196597 BM196597
33	8	40.0	490	9	AV890393	AV890393 AV890393
34	8	40.0	493	9	AV862905	AV862905 AV862905
C 35	8	40.0	501	13	BM202753	BM202753 BM202753
C 36	8	40.0	504	14	CD112744	CD112744 ME1-0024T
C 37	8	40.0	505	14	CA820687	CA820687 EST0012 N
C 38	8	40.0	510	9	AV950469	AV950469 AV950469
C 39	8	40.0	510	29	TA90C02P	TA90C02P T. brucei
C 40	8	40.0	515	29	CC476294	CC476294 CH240_302
C 41	8	40.0	516	12	BP001077	BP001077 BP001077
C 42	8	40.0	521	13	BW136624	BW136624 BW136624
C 43	8	40.0	524	13	BW203262	BW203262 BW203262
C 44	8	40.0	525	9	AV680124	AV680124 AV680124
C 45	8	40.0	526	9	AV973566	AV973566 AV973566

# ALIGNMENTS

RESULT 1  
BF910533/c  
LOCUS  
DEFINITION  
CM4-UT0042-011100-395-f08 UT0042 Homo sapiens cDNA, mRNA sequence.  
EST 18-JAN-2001  
ACCESSION  
BF910533  
VERSION  
BF910533.1 GI:12301991  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens (human)  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE  
1 (bases 1 to 240)  
Dias Neto,E., Garcia Correa,R., Verjovskij-Almeida,S., Briones,M.R.,

Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deOliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.

Shotgun sequencing of the human transcriptome with ORF expressed

sequence tags

JOURNAL

MEDLINE

PUBMED

COMMENT

Contact: Simpson A.J.G.

Laboratory of Cancer Genetics

Ludwig Institute for Cancer Research

Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil

Tel: +55-11-2704922

Fax: +55-11-2707001

Email: asimpson@ludwig.org.br

This sequence was derived from the FAPESP/LICR Human Cancer Genome

Project. This entry can be seen in the following URL

(<http://www.ludwig.org.br/scripts/gethtml2.pl?tl=CM4&t2=CM4-UT0042-011100-395-f08&t3=2000-11-01&t4=1>)

Seq primer: puc 18 forward

High quality sequence start: 81

High quality sequence stop: 240.

Location/Qualifiers

FEATURES

source

1..240

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/dev\_stage="Adult"

/clone\_lib="UT0042"

/note="Organ: uterus\_tumor; Vector: puc18; Site\_1: Smar;

Site\_2: Smal; A mini-library was made by cloning products

derived from ORESTES PCR (U.S. Letters Patent application

No. 196,716 - Ludwig Institute for Cancer Research)

profiles into the puc 18 vector. Reverse transcription of

tissue mRNA and cDNA amplification were performed under

low stringency conditions."

ORIGIN

Alignment Scores:  
Pred. No.: 6,33e-09 Length: 240  
Score: 19.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 95.00% Indels: 0  
DB: 10 Gaps: 0

US-10-799-747-116 (1-20) x BF910533 (1-240)

Qy 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyAlaLeuLysSer 19

Db 191 ATGGCAGCCCAATTCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTCTGAAATCA 135

RESULT 2

BM722991

LOCUS

DEFINITION UI-E-EO1-aid-a-14-0-UI.r1 UI-E-EO1 Homo sapiens cDNA clone

UI-E-EO1-aid-a-14-0-UI 5', mRNA sequence.

ACCESSION BM722991

VERSION BM722991.1 GI:19043963

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 575)

Bonaldo,M.F., Lennon,G. and Soares,M.B.

Normalization and subtraction: two approaches to facilitate gene

discovery

Genome Res. 6 (9), 791-806 (1996)

JOURNAL

MEDLINE

97044477

PUBMED

COMMENT

8889548

Contact: Soares, MB

Coordinated Laboratory for Computational Genomics

University of Iowa

375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA

Tel: 319 335 8250

Fax: 319 335 9565

Email: bento-soares@uiowa.edu

Tissue Procurement: Dr. Gregg Hageman

cDNA Library preparation: Dr. M. Bento Soares, University of Iowa

cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa

Clone Distribution: Researchers may obtain clones from Research

Genetics ([www.resgen.com](http://www.resgen.com)).

The following repetitive elements were found in this cDNA

sequence: 148-169, >AT-rich#Low\_complexity (matched complement)

Seq primer: M13 Reverse.

Location/Qualifiers

source

1..575

/organism="Homo sapiens"

/mol\_type="mRNA"

/db\_xref="taxon:9606"

/clone="UI-E-EO1-aid-a-14-0-UI"

/tissue\_type="fetal eye"

/dev\_stage="fetal"

/lab\_host="DH10B (Life Technologies) (T1 phage resistant)"

/clone\_lib="UI-E-EO1"

/note="Organ: eye; Vector: pT73-Pac (Pharmacia) with a

modified polylinker; Site\_1: EcoR I; Site\_2: Not I;

UI-E-EO1 is a normalized cDNA library containing the

following tissue(s): fetal eye. The library was

constructed according to Bonaldo, Lennon and Soares,

Genome Research, 6:791-806, 1996. First strand cDNA

synthesis was primed with an oligo-dT primer containing a

Not I site. Double stranded cDNA was ligated to an EcoR I

adaptor, digested with Not I, and cloned directionally

into pT73-Pac vector. The oligonucleotide used to prime

the synthesis of first-strand cDNA contains a library tag

sequence that is located between the Not I site and the

(dT)18 tail. The sequence tag for this library is

CGCGTATACC. This library was created for the program, Gene

Discovery in the Visual System, supported by National Eye

Institute (NEI)."

ORIGIN

Alignment Scores:  
Pred. No.: 1.55e-08 Length: 575  
Score: 19.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 95.00% Indels: 0  
DB: 12 Gaps: 0

US-10-799-747-116 (1-20) x BM722991 (1-575)

Qy 1 MetAlaAlaHisSerValLeuSerPheLeuLeuThrProTyAlaLeuLysSer 19

Db 454 ATGGCAGCCCAATTCAGTCTTGAGTTTCTCTCTGGACACCTTATGCTCTGAAATCA 510

RESULT 3

BI914473

LOCUS

DEFINITION BI914473 NIH\_MGC\_121 Homo sapiens cDNA clone IMAGE:5246463 5',

mRNA sequence.

ACCESSION BI914473

VERSION BI914473.1 GI:16178652

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 765)

NIH-MGC <http://mgc.nci.nih.gov/>.

REFERENCE

AUTHORS

**TITLE** National Institutes of Health, Mammalian Gene Collection (MGC)  
**JOURNAL** Unpublished (1999)  
**COMMENT** Contact: Robert Strausberg, Ph.D.  
 Email: cgapbs-r@mail.nih.gov  
 Tissue Procurement: Life Technologies, Inc.  
 cDNA Library Preparation: Life Technologies, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
 Plate: LLAM11621 row: p column: 16  
 High quality sequence stop: 762.

**FEATURES** Location/Qualifiers  
 source  
 1..765  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:5246463"  
 /lab\_host="DH10B"  
 /clone\_lib="NIH\_MGC\_121"  
 /note="Organ: brain; Vector: pCMV-SPORT6; Site 1: NotI; Site 2: EcoRV (destroyed); RNA source anonymous pool of 3 fetal brains, female age 20 weeks, female age 24 weeks, and male age 26 weeks. Library is oligo-dT primed and directionally cloned (EcoRV site is destroyed upon cloning). Average insert size 1.7 kb, insert size range 0.7-3.5 kb. Library is normalized and enriched for full-length clones and was constructed by C. Gruber (Invitrogen). Research Genetics tracking code 017. Note: this is a NIH\_MGC Library."

**Alignment Scores:**  
 Pred. No.: 2.07e-08 Length: 765  
 Score: 19.00 Matches: 19  
 Percent Similarity: 100.00% Conservatives: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 95.00% Indels: 0  
 DB: 12 Gaps: 0

US-10-799-747-116 (1-20) x BI914473 (1-765)

**Qy** 1 MetaAlaHisSerValLeuSerPheLeuLeuTrpThrProTyAlaLeuLysSer 19  
 |||||  
 Db 556 ATGCAGCCCATTCAGTTGAGTTTCTCTCTGGACACCTATGCTCTGAATCA 612

**RESULT 4**  
**LOCUS** BG484396 785 bp mRNA linear EST 21-MAR-2001  
**DEFINITION** 602505037F1 NIH\_MGC\_77 Homo sapiens cDNA clone IMAGE:4618473 5', mRNA sequence.  
**ACCESSION** BG484396  
**VERSION** BG484396.1 GI:13416675  
**KEYWORDS** EST.  
**SOURCE** Homo sapiens (human)  
**ORGANISM** Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 785)  
**REFERENCE** NIH-MGC <http://mgc.nci.nih.gov/>.  
**AUTHORS** National Institutes of Health, Mammalian Gene Collection (MGC)  
**TITLE** Unpublished (1999)  
**JOURNAL** Contact: Robert Strausberg, Ph.D.  
**COMMENT** Email: cgapbs-r@mail.nih.gov  
 Tissue Procurement: CLONTECH Laboratories, Inc.  
 cDNA Library Preparation: CLONTECH Laboratories, Inc.  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
 Plate: LLCM1376 row: j column: 10

High quality sequence start: 18  
 High quality sequence stop: 751.

**FEATURES** Location/Qualifiers  
 source  
 1..785  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="IMAGE:4618473"  
 /lab\_host="DH10B (T1 phage-resistant)"  
 /clone\_lib="NIH\_MGC\_77"  
 /note="Organ: lung; Vector: pDNR-LIB (Clontech); Site 1: SfiI (ggccgtctggcc); Site 2: SfiI (ggccattggcc); 5' and 3' adaptors were used in cloning as follows: 5' adaptor sequence: 5'-CACGCCATTATGCC-3' and 3' adaptor sequence: 5'-ATTCTAGAGCGCGCGCGACATG-dT(30)BN-3' (where B = A, C, or G and N = A, C, G, or T). Average insert size 1.9 kb (range 0.5-4.0 kb). 12/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA). Note: this is a NIH\_MGC Library."

**ORIGIN**  
 Alignment Scores:  
 Pred. No.: 2.13e-08 Length: 785  
 Score: 19.00 Matches: 19  
 Percent Similarity: 100.00% Conservatives: 0  
 Best Local Similarity: 100.00% Mismatches: 0  
 Query Match: 95.00% Indels: 0  
 DB: 12 Gaps: 0

US-10-799-747-116 (1-20) x BG484396 (1-785)

**Qy** 1 MetaAlaHisSerValLeuSerPheLeuLeuTrpThrProTyAlaLeuLysSer 19  
 |||||  
 Db 322 ATGCAGCCCATTCAGTTGAGTTTCTCTCTGGACACCTATGCTCTGAATCA 378

**RESULT 5**  
**LOCUS** BX441923 1201 bp mRNA linear EST 15-MAY-2003  
**DEFINITION** BX441923 Homo sapiens FETAL BRAIN Homo sapiens cDNA clone CSODF023Y002 5-PRIME, mRNA sequence.  
**ACCESSION** BX441923  
**VERSION** BX441923.1 GI:30771989  
**KEYWORDS** EST.  
**SOURCE** Homo sapiens (human)  
**ORGANISM** Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 1201)  
**REFERENCE** Li, W.B., Gruber, C., Jesse, J. and Polayes, D.  
**AUTHORS** Full-length cDNA libraries and normalization  
**TITLE** Unpublished (2001)  
**JOURNAL** Contact: Genoscope  
**COMMENT** Genoscope - Centre National de Sequencage  
 BP 191 91006 EVRY cedex - France  
 Email: seqref@genoscope.cns.fr, Web : [www.genoscope.cns.fr](http://www.genoscope.cns.fr)  
 Library was constructed by Life Technologies, a division of Invitrogen. This sequence belongs to sequence cluster 2532.r For more information about this cluster, see <http://www.genoscope.cns.fr/cgi-bin/cluster.cgi?seq=CSODF023BH01QPI&cluster=2532.r>. Contact : Feng Liang Email : [fliang@lifetech.com](mailto:fliang@lifetech.com) URL : <http://fulllength.invitrogen.com/> Invitrogen Corporation 1600 Faraday Avenue Genoscope sequence ID : CSODF023BH01QPI.

**FEATURES** Location/Qualifiers  
 source  
 1..1201  
 /organism="Homo sapiens"  
 /mol\_type="mRNA"  
 /db\_xref="taxon:9606"  
 /clone="CSODF023Y002"  
 /tissue\_type="FETAL BRAIN"  
 /dev\_stage="Fetal"  
 /clone\_lib="Homo sapiens FETAL BRAIN"

/note="Organ: brain; Vector: pCMVSPORT 6; 1st strand cDNA was primed with a NotI-oligo(dT) primer. Five prime end enriched, double-strand cDNA was digested with Not I and cloned into the Not I and EcoRV sites of the pCMVSPORT 6 vector. Library was not normalized."

## ORIGIN

Alignment Scores:  
Pred. No.: 3.28e-08 Length: 1201  
Score: 19.00 Matches: 19  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 95.00% Indels: 0  
DB: 13 Gaps: 0

US-10-799-747-116 (1-20) x BX441923 (1-1201)

QY 1 MetAlaAlaHisSerValLeuSerPheLeuLeuTTPThrProTyAlaLeuLySer 19

Db 146 ATGGCAGCCCATTCAGTCTTGAGTTTCTTCTGACACCTTATGCTCGAATCA 202

## RESULT 6

AL035941 621 bp mRNA linear EST 04-SEP-2003  
LOCUS DKZP564B1622 r1.564 (synonym: hfr2) Homo sapiens cDNA clone  
DEFINITION DKZP564B1622 5', mRNA sequence.

ACCESSION AL035941

VERSION AL035941

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 621)

AUTHORS Wambutt, R., Heubner, D., Mewes, H.W., Gassenhuber, J. and Wiemann, S.

TITLE EST (Wambutt, et al.)

JOURNAL Unpublished (1999)

COMMENT Contact: MIPS

MIPS

Ingolstaedter Landstr.1, D-85764 Neuherberg, Germany

This is the 5' sequence of the clone insert

Clone from S. Wiemann, Molecular Genome Analysis, German Cancer

Research Center (DKFZ); Email: s.wiemann@dkfz-heidelberg.de;

sequenced by AGOWA (Berlin/Germany) within the cDNA sequencing

consortium of the German Genome Project.

s1 sequence also available.

This clone (DKFZP564B1622) is available at the RZPD in Berlin.

Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059

Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

## FEATURES

source

1..621  
Location/Qualifiers  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="taxon:9606"  
/clone="DKFZP564B1622"  
/tissue\_type="brain"  
/dev\_stage="fetal"  
/lab\_host="xl-2blue"  
/clone\_lib="564 (synonym: hfr2)"  
/note="Vector: pAMP1; Site\_1: NotI; Site\_2: SalI"

## ORIGIN

Alignment Scores:  
Pred. No.: 6.57 Length: 621  
Score: 10.00 Matches: 10  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 50.00% Indels: 0  
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AL035941 (1-621)

QY 1 MetAlaAlaHisSerValLeuSerPheLeu 10

Db 591 ATGGCAGCCCATTCAGTCTTGAGTTTCTT 620

## RESULT 7

LOCUS AQ293004

DEFINITION

HS\_2225\_A1\_E02\_MF CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=2225 Col=3 Row=I, genomic survey sequence.

ACCESSION AQ293004

VERSION AQ293004.1

KEYWORDS GSS.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 496)

AUTHORS Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and Hood, L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

MEDLINE 99380589

PUBMED 10449764

COMMENT Contact: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Place: 2225 row: I column: 3

Class: BAC ends

High quality sequence stop: 496.

Location/Qualifiers

1..496

/organism="Homo sapiens"

/mol\_type="genomic DNA"

/db\_xref="taxon:9606"

/clone="Plate=2225 Col=3 Row=I"

/sex="male"

/clone\_lib="CIT Approved Human Genomic Sperm Library D"

/note="Organ: sperm; Vector: pBelobAC11; BAC Clones in E-Coli DH10B"

ORIGIN

Alignment Scores:

Pred. No.: 47.1 Length: 496

Score: 9.00 Matches: 9

Percent Similarity: 100.00% Conservative: 0

Best Local Similarity: 100.00% Mismatches: 0

Query Match: 45.00% Indels: 0

DB: 28 Gaps: 0

US-10-799-747-116 (1-20) x AQ293004 (1-496)

QY 5 SerValLeuSerPheLeuLeuTrpThr 13

Db 119 TCTGTATTATCTCTCTGCTATGACT 145

## RESULT 8

LOCUS AV851019

DEFINITION

AV851019 Nori Satoh unpublished cDNA library, larva Clona intestinalis cDNA clone rcilv10n02 3', mRNA sequence.

ACCESSION AV851019

VERSION AV851019.1

KEYWORDS EST.

SOURCE Clona intestinalis

ORGANISM Clona intestinalis

Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;

AV851019 239 bp mRNA linear EST 08-NOV-2001

AV851019 Nori Satoh unpublished cDNA library, larva Clona

intestinalis cDNA clone rcilv10n02 3', mRNA sequence.

AV851019

AV851019.1 GI:16834788

EST.

Clona intestinalis

Clona intestinalis

Eukaryota; Metazoa; Chordata; Urochordata; Ascidiacea; Enterogona;

REFERENCE  
1 (bases 1 to 239)  
Satcho, N., Satcho, Y., Kohara, Y. and Shin-i, T.  
Expressed genes in Ciona intestinalis  
JOURNAL  
COMMENT  
Contact: Nori Satcho  
Department of Zoology  
Kyoto University  
Sakyo-ku, Kyoto, Kyoto 606-8502, Japan  
Tel: 81-75-753-4081  
Fax: 81-75-705-1113  
Email: satcho@ascidian.zool.kyoto-u.ac.jp.

FEATURES  
source  
Location/Qualifiers

1..239  
/organism="Ciona intestinalis"  
/mol\_type="mRNA"  
/db\_xref="taxon:7719"  
/clone="rcilv10n02"  
/tissue\_type="whole animal"  
/dev\_stage="larva"  
/clone\_lib="Nori Satcho unpublished cDNA library, larva"

ORIGIN

Alignment Scores:  
Pred. No.: 201 Length: 239  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 9 Gaps: 0

US-10-799-747-116 (1-20) x AV851019 (1-239)

Qy 4 HisServValLeuSerPheLeu 11  
Db 173 CATTGAGTTTGTCTTTTTCCTC 196

RESULT 9  
BG371993  
LOCUS  
DEFINITION  
UI-R-CVO-brl-a-01-0-UI-s1 UI-R-CVO Rattus norvegicus cDNA clone  
UI-R-CVO-brl-a-01-0-UI 3', mRNA sequence.  
ACCESSION  
BG371993  
VERSION  
BG371993.1 GI:13268530  
KEYWORDS  
EST.  
SOURCE  
Rattus norvegicus (Norway rat)  
ORGANISM  
Rattus norvegicus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;  
Rattus.

REFERENCE  
AUTHORS  
TITLE  
Normalization and subtraction: two approaches to facilitate gene  
discovery  
JOURNAL  
Genome Res. 6 (9), 791-806 (1996)  
MEDLINE  
97044477  
PUBMED  
8889548

COMMENT  
Contact: Soares, MB  
Coordinated Laboratory for Computational Genomics  
University of Iowa  
375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA  
Tel: 319 335 8250  
Fax: 319 335 9565  
Email: bento-soares@uiowa.edu

The sequence contained an oligo-dT track that was present in the  
oligonucleotide that was used to prime the synthesis of first  
strand cDNA and therefore this may represent a bonafide poly A  
tail. The sequence tag present in the cDNA between the NotI site  
and the oligo-dT track served to verify it as a clone from the  
non-normalized rat eye library cDNA Library Preparation: M.B.  
Soares Lab Clone distribution: clones will be available through  
Research Genetics (www.resgen.com) The following repetitive  
elements were found in this cDNA sequence: 1-25,

>AT rich#Low complexity  
Seq primer: M13 Forward  
POLYA=Yes.

FEATURES  
source  
Location/Qualifiers

1..272  
/organism="Rattus norvegicus"  
/mol\_type="mRNA"  
/strain="Sprague-Dawley"  
/db\_xref="taxon:10116"  
/clone="UI-R-CVO-brl-a-01-0-UI"  
/dev\_stage="ADULT"  
/lab\_host="DH10B (Life Technologies)"  
/clone\_lib="UI-R-CVO"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified  
polylinker; Site 1: Not I; Site 2: Eco RI; The UI-R-CVO  
library is a non-normalized library constructed from rat  
eye tissue. For a detailed description of the library from  
which this clone was derived, please visit our web site at  
rategest.eng.uiowa.edu. The subtraction has been previously  
described in (Bonaldo, Lennon and Soares, Genome Research  
6:791-806, 1996)  
TAG\_TISSUE=rat eye  
TAG\_LIB=UI-R-CVO  
TAG\_SEQ=CAGCC"

ORIGIN

Alignment Scores:  
Pred. No.: 230 Length: 272  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 12 Gaps: 0

US-10-799-747-116 (1-20) x BG371993 (1-272)

Qy 2 AlaAlaHisServValLeuSerPhe 9  
Db 107 GCAGCACATTCGTGCTTCTTC 130

RESULT 10  
CE488576/c  
LOCUS  
DEFINITION  
tigr-gss-dog-17000365567544 Dog Library Canis familiaris genomic,  
genomic survey sequence.  
ACCESSION  
CE488576  
VERSION  
CE488576.1 GI:36805357  
KEYWORDS  
GSS.  
SOURCE  
Canis familiaris (dog)

ORGANISM  
Canis familiaris  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.

REFERENCE  
AUTHORS  
1 (bases 1 to 290)  
Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,  
Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and  
Venter, J.C.

TITLE  
The dog genome: survey sequencing and comparative analysis  
JOURNAL  
Science 301 (5641), 1898-1903 (2003)  
MEDLINE  
22875432  
PUBMED  
14512627

COMMENT  
Contact: Kirkness EF  
The Institute for Genomic Research  
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,  
Rockville, MD 20850, USA  
Tel: 301-838-0200  
Fax: 301-838-0208  
Email: ekirknes@tigr.org  
Class: shotgun.

FEATURES  
source  
Location/Qualifiers

1..290  
/organism="Canis familiaris"  
/mol\_type="genomic DNA"  
/strain="Standard Poodle"



ORGANISM Schistosoma mansoni  
Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigeida; Schistosomatoidea; Schistosomatidae; Schistosoma.  
1 (bases 1 to 336)

REFERENCE  
AUTHORS Verjovski-Almeida, S., DeMarco, R., Martins, E.A.L., Guimaraes, P.E.M.,  
Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,  
Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,  
Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,  
Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A.,  
Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,  
Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,  
Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,  
Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.

TITLE Transcriptome analysis of the acelomate human parasite Schistosoma  
mansoni

JOURNAL Nat. Genet. 35 (2), 148-157 (2003)  
MEDLINE 22879926  
COMMENT Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP,  
Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjo@iq.usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
Genome Project. All sequences in the project were assembled and  
annotated. This entry and all the assembled sequences can be seen  
in the following URL <http://bioinfo.iq.usp.br/schisto/>  
Plate: MSI-0139P-V386 row: 1 column: G.

FEATURES  
Location/Qualifiers  
source  
1..336  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MSI-0139P-V386-G01.B"  
/sex="mixed pool"  
/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_lib="MSI-0139"  
/note="Vector: pGEM T-easy"

ORIGIN  
Alignment Scores:  
Pred. No.: 285 Length: 336  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 14 Gaps: 0

US-10-799-747-116 (1-20) x CD202965 (1-336)

Qy 4 HisSerValLeuSerPheLeu 11  
|||||  
Db 99 CATTCGGTTTAAAGTTTCTCTG 122  
|||||

RESULT 14  
T59368  
LOCUS yb57h01.sl Stragatene ovary (#937217) Homo sapiens cDNA clone  
DEFINITION IMAGE:7513 3' similar to gb:X77738 rnal BAND 3 ANION TRANSPORT  
PROTEIN (HUMAN); contains Alu repetitive element; mRNA sequence.  
T59368  
ACCESSION T59368.1 GI:661205  
VERSION T59368.1  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE  
AUTHORS 1 (bases 1 to 356)  
Hallier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiappelli, B.,  
Chissoe, S., Dietrich, N., DuBuque, T., Favello, A., Gish, W.,

ORGANISM Schistosoma mansoni  
Eukaryota; Metazoa; Platyhelminthes; Trematoda; Digenea;  
Strigeida; Schistosomatoidea; Schistosomatidae; Schistosoma.  
1 (bases 1 to 336)

REFERENCE  
AUTHORS Verjovski-Almeida, S., DeMarco, R., Martins, E.A.L., Guimaraes, P.E.M.,  
Ojopi, E.P.B., Paquola, A.C.M., Piazza, J.P., Nishiyama, M.Y. Jr.,  
Kitajima, J.P., Adamson, R.E., Ashton, P.D., Bonaldo, M.F.,  
Coulson, P.S., Dillon, G.P., Farias, L.P., Gregorio, S.P., Ho, P.L.,  
Leite, R.A., Malaquias, L.C.C., Marques, R.C.P., Miyasato, P.A.,  
Nascimento, A.L.T.O., Ohlweiler, F.P., Reis, E.M., Ribeiro, M.A.,  
Sa, R.G., Stukart, G.C., Soares, M.B., Gargioni, C., Kawano, T.,  
Rodrigues, V., Madeira, A.M.B.N., Wilson, R.A., Menck, C.F.M.,  
Setubal, J.C., Leite, L.C.C. and Dias-Neto, E.

TITLE Transcriptome analysis of the acelomate human parasite Schistosoma  
mansoni

JOURNAL Nat. Genet. 35 (2), 148-157 (2003)  
MEDLINE 22879926  
COMMENT Contact: Dr. Sergio Verjovski-Almeida  
Departamento de Bioquímica  
Instituto de Química - Universidade de São Paulo  
Av. Prof. Lineu Prestes 748 sala 1200, 05508-900 São Paulo - SP,  
Brasil  
Tel: +55-11-3091-2173  
Fax: +55-11-3091-2186  
Email: verjo@iq.usp.br  
This sequence was derived from the FAPESP Schistosoma mansoni EST  
Genome Project. All sequences in the project were assembled and  
annotated. This entry and all the assembled sequences can be seen  
in the following URL <http://bioinfo.iq.usp.br/schisto/>  
Plate: MSI-0139P-V386 row: 1 column: G.

FEATURES  
Location/Qualifiers  
source  
1..336  
/organism="Schistosoma mansoni"  
/mol\_type="mRNA"  
/db\_xref="taxon:6183"  
/clone="MSI-0139P-V386-G01.B"  
/sex="mixed pool"  
/dev\_stage="schistosomulum"  
/lab\_host="in vitro culture"  
/clone\_lib="MSI-0139"  
/note="Vector: pGEM T-easy"

ORIGIN  
Alignment Scores:  
Pred. No.: 285 Length: 336  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 14 Gaps: 0

US-10-799-747-116 (1-20) x CD202965 (1-336)

Qy 4 HisSerValLeuSerPheLeu 11  
|||||  
Db 99 CATTCGGTTTAAAGTTTCTCTG 122  
|||||

RESULT 14  
T59368  
LOCUS yb57h01.sl Stragatene ovary (#937217) Homo sapiens cDNA clone  
DEFINITION IMAGE:7513 3' similar to gb:X77738 rnal BAND 3 ANION TRANSPORT  
PROTEIN (HUMAN); contains Alu repetitive element; mRNA sequence.  
T59368  
ACCESSION T59368.1 GI:661205  
VERSION T59368.1  
KEYWORDS EST.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE  
AUTHORS 1 (bases 1 to 356)  
Hallier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiappelli, B.,  
Chissoe, S., Dietrich, N., DuBuque, T., Favello, A., Gish, W.,

Hawkins, M., Hultman, M., Kucaba, T., Lacy, M., Le, N.,  
Mardis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L.,  
Rohlfing, T., Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J.,  
Trevaskis, E., Underwood, K., Wohlmann, P., Waterston, R., Wilson, R.  
and Marra, M.  
Generation and analysis of 280,000 human expressed sequence tags  
Genome Res. 6 (9), 807-828 (1996)  
97044478  
8889549  
PUBMED  
COMMENT  
Contact: Wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: est@wustl.wustl.edu  
High quality sequence stops: 339  
Source: IMAGE Consortium, LINL  
This clone is available royalty-free through LINL; contact the  
IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.  
Seq primer: -21ml3  
High quality sequence stop: 339.  
Location/Qualifiers  
source  
1..356  
/organism="Homo sapiens"  
/mol\_type="mRNA"  
/db\_xref="GDB:505042"  
/db\_xref="taxon:9606"  
/clone="IMAGE:75313"  
/sex="female"  
/dev\_stage="49 year old"  
/lab\_host="SOLR cells (kanamycin resistant)"  
/clone\_lib="Stratagene ovary (#937217)"  
/note="Organ: ovary; Vector: Bluescript SK; Site 1: EcoRI;  
Site 2: XhoI; Cloned unidirectionally. Primer: Oligo dT.  
Total ovary tissue, normal, caucasian. Average insert  
size: 0.8 Kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5'  
GAATTCGGCAGG 3' -3' adaptor sequence: 5'  
CTCGAGTCTTTTCTTTTCTTTT 3'"

ORIGIN  
Alignment Scores:  
Pred. No.: 302 Length: 356  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 14 Gaps: 0

US-10-799-747-116 (1-20) x T59368 (1-356)

Qy 3 AlahSerValLeuSerPheLeu 10  
|||||  
Db 108 GTCATTCGTGACTTCTTCTCTC 131  
|||||

RESULT 15  
CE286752  
LOCUS tigr-gss-dog-1700033810288 Dog Library Canis familiaris genomic,  
DEFINITION genomic survey sequence.  
ACCESSION CE286752  
VERSION CE286752.1 GI:36056365  
KEYWORDS GSS.  
SOURCE Canis familiaris (dog)  
ORGANISM Canis familiaris  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Carnivora; Fissipedia; Canidae; Canis.  
REFERENCE  
AUTHORS 1 (bases 1 to 385)  
Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,  
Rusch, D.B., Deicher, A.L., Pop, M., Wang, W., Fraser, C.M. and  
Venter, J.C.  
The dog genome: survey sequencing and comparative analysis  
Science 301 (5641), 1898-1903 (2003)  
MEDLINE 22875432

PUBMED 14512627  
COMMENT Contact: Kirkness EF  
The Institute for Genomic Research  
Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,  
Rockville, MD 20850, USA  
Tel: 301-838-0200  
Fax: 301-838-0208  
Email: ekirknes@tigr.org  
Class: shotgun.  
FEATURES Location/Qualifiers  
source 1..385  
/organism="Canis familiaris"  
/mol\_type="genomic DNA"  
/strain="Standard Poodle"  
/db\_xref="taxon:9615"  
/clone\_lib="Dog Library"  
/note="Site 1: BstXI; Libraries were prepared from  
peripheral blood"

ORIGIN

Alignment Scores:  
Pred. No.: 327 Length: 385  
Score: 8.00 Matches: 8  
Percent Similarity: 100.00% Conservative: 0  
Best Local Similarity: 100.00% Mismatches: 0  
Query Match: 40.00% Indels: 0  
DB: 29 Gaps: 0

US-10-799-747-116 (1-20) x CE286752 (1-385)

QY 4 HisSerValLeuSerPheLeuLeu 11  
DB 198 CACAGCGTCTATCAATTCCTCTG 221

Search completed: July 21, 2004, 05:22:32  
Job time : 2475 secs